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AN ASSAY DEVICE

Abstract:

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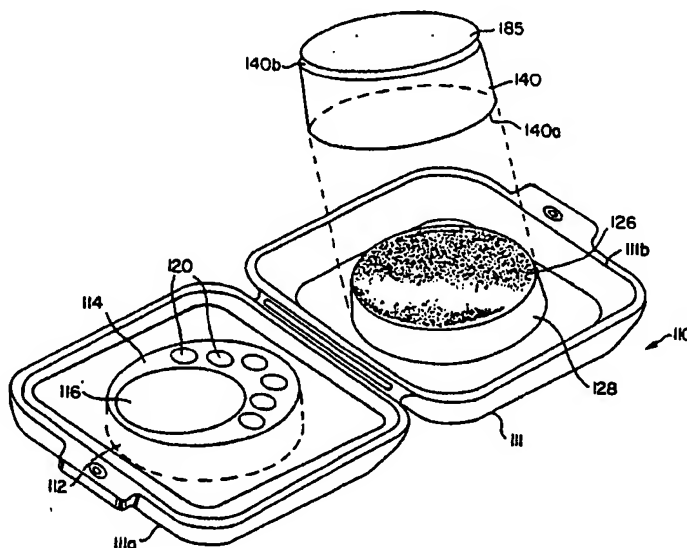
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(54) Title: AN ASSAY DEVICE



(57) Abstract

An assay device and method are provided which allow for determination of the presence of an analyte in a sample while providing identification of the test subject. Self-contained assay device (110) includes a housing (111) with a first part (111a) and a second part (111b). Reaction medium (114) is located within the first part (111a) and mounted on a base member (112). Base member (112) includes a porous support which may be adjacent to the reaction medium (114) which includes at least reaction zone (120) as well as a control zone (116). A medium (126) containing a signal-producing agent is located in the second part of the housing (111b) and mounted on a base portion (128). The device additionally comprises a cylindrical body (140) having an open end (140a) and a closed end (140b) on which is mounted a medium (185) for containing and transferring a portion of test sample to the reaction medium (114).

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AN ASSAY DEVICE

This application is a continuation-in-part application of International Application No. PCT/US92/07785, filed on September 14, 1992 and of a
5 continuation application filed on September 9, 1993, but which has not yet received a serial no. of United States Serial No. 07/759,922, filed on September 13, 1991 (and from which PCT/US92/07785
10 claims priority), which is a continuation-in-part application of United States Application Serial No. 07/668,115, filed on March 12, 1991 (now abandoned), which is a continuation-in-part application of United States Patent Application Serial No. 07/467,532, filed on January 19, 1990 (which is now
15 abandoned), all of which are incorporated herein by reference.

TECHNICAL FIELD OF THE INVENTION

The invention is in the field of ligand-receptor assays, including immunoassays, used
20 for determining the presence or absence of an analyte in a biological fluid. More particularly, the apparatuses and methods of the invention relate to establishing the identity of a test subject and determining the presence or absence of at least one
25 analyte in a biological fluid taken from the test subject using a single test device.

BACKGROUND OF INVENTION

For many years, those skilled in the art of ligand-receptor assays have sought an effective,
30 inexpensive, and reliable device and method for detecting the presence and/or absence of antigens,

antibodies, and the like. The art is replete with such devices.

For many years, those skilled in the art have also sought devices and methods for assuring that
5 the sample being tested actually came from a certain individual. Many schemes, some elaborate, some simple, have been established to verify that the test sample was in fact produced by a particular individual. Similarly, many schemes exist to thwart
10 matching a particular test result with a particular person.

For example, drug testing is now a routine procedure in athletics, prison, and the work place. The protocols involve testing individual fluid
15 samples such as urine or blood to determine the presence of certain antibodies in the fluid sample; a positive result may be an indication of drug use.

Previously such testing has been accomplished by a series of tests which may involve shifting of
20 the fluid being tested to different containers and removal of the fluid from the person being tested to a distant place. Oftentimes, fluid misplacement and/or substitution opened questions as to the chain of custody of the tested fluid. Thus, a problem
25 which has occurred during such testing is that test fluids may be obtained from persons other than the person to be tested or that test fluids become mixed, lost, or cannot be specifically identified with that person after the test is returned from the
30 laboratory. Also, these assays typically take too long to obtain timely results.

Other shortcomings are associated with many of

the devices and methods currently used for the determination of analytes in biological fluids. Frequently, a certain level of skill is required to use or handle such devices to obtain satisfactory results. In addition, most of these devices require one or more additional reagents, and frequently a wash solution, to apply to the device during the course of the test. In many of these devices, due at least in part to the limited sensitivity of the devices, relatively large amounts of oftentimes expensive reagents or of test sample are required to give accurate results.

As noted above, the principles which form the basis for this type of detection is a person's immune system, i.e., the inherent capability of a mammal to respond to a foreign molecule, typically a macromolecule. Hereinafter, any molecule which is capable of eliciting such a response will be referred to as an antigen, i.e., an antibody generator. The proteins and protein fragments which are produced in response to the antigen will be referred to as antibodies or immunoglobulins.

SUMMARY OF THE INVENTION

The present invention overcomes problems inherent in known assay devices and systems through the use of a specifically designed assay device, preferably an immunoassay device which allows the determination of the presence or absence of an analyte in a sample, and preferably specifically identifies the person providing the sample.

The present invention is directed to a testing

method and device and more specifically to a method and device for detecting the presence of an analyte, most preferably specific antigens or specific antibodies in a biological fluid. The present invention preferably also provides for positively identifying the individual tested.

The present invention provides an easily handled, disposable testing pad and a disposable test device suitable for detecting the presence or absence of an analyte and preferably for identifying the test subject. For example, a finger of the test subject may be coated with a labelled ligand and the finger may be pressed on a membrane substrate having a specific immobilized ligand bed to capture an analyte from a biological fluid while simultaneously providing an inkless (specific binding) visible fingerprint of the test subject so that positive identification of the fluid donor is irrefutably obtained. The fingerprint of the fluid donor obtained on the membrane may also include information about analytes, such as drugs, present in his body fluids at the time of performing the test. This information can be recorded or stored for positive identification when needed.

It is an object of the invention to assay a test sample for the presence or absence of an analyte while also identifying the subject being tested. Thus, problems of misidentification, chain of title, and delays may be eliminated.

It is an object of the present invention to provide a device for rapidly determining the presence or absence of an analyte taken from a test subject. It is also an object of the present

invention to provide a device for rapidly determining the quantitative amount of an analyte in a sample taken from a test subject. Yet other objects of the present invention are to provide a device and a method of determining the presence or absence of a drug or drug metabolite in a test sample taken from a test subject and to provide a device and method determining whether a test subject is an acute or chronic drug abuser.

Another embodiment of the present invention, where identifying the test subject by a fingerprint is not critical, involves applying the sample of body fluid or a labelled ligand to the membrane with the pressure of a transfer means or applicator that is part of the test device. In contrast to similar known devices which apply liquid test samples and/or reagents dropwise to the surface of a reaction medium, the present invention, in which liquid test samples and/or reagents are applied and maintained under pressure on the surface of the reaction medium for a short duration, requires smaller amounts of test sample and reagents and demonstrates higher sensitivities than known devices.

A device according to the present invention which uses this principle is an enclosed assay test device. A preferred embodiment is one in which the assay test device includes a housing, a reaction medium having at least one reaction zone located in the housing, a medium containing a signal-producing or analyte-indicating agent located in the housing, and a medium for containing a test sample located in the housing. Both the medium containing a signal-producing agent and the medium for containing

a test sample located in the housing are movable independently between a first position in spaced relationship to the reaction medium and a second position in substance-transferrable contact with the reaction medium. Preferably this is a self-contained test device.

A particular embodiment of this invention includes an assay test device employing a two-part housing. In one embodiment of the device, a reaction medium having at least one reaction zone is mounted in a first part of the housing. In the second part of the two-part housing is located a medium for containing a portion of a test sample. The medium for containing a portion of a test sample and the reaction medium are so arranged in the two parts of the housing that when the housing parts are brought together in a closed relationship, there is substance transferrable contact between the medium for containing a portion of a test sample and the reaction medium. In addition to at least one reaction zone, the reaction medium preferably has a control zone which is provided to identify the test subject by a fingerprint or like means if desired. Such a device may be designed as a self-contained assay device. As used herein, the term "self-contained" means that the device may be used without the addition of additional reagents or solutions, or only requiring the dropwise addition of water.

Another embodiment employs a somewhat similar arrangement but a medium containing a signal-producing agent is substituted for the medium for containing a portion of a test sample.

A preferred device according to the above

"pressure" principle is similar to the
aforementioned devices in that a two-part housing is
also employed in which a reaction medium having at
least one reaction zone is mounted in a first part
5 of the two-part housing. However, like one of these
embodiments, in the second part of the two-part
housing, there is located a medium containing a
signal-producing agent, such as a labelled material
which will indicate the presence of an analyte. The
10 device also includes a removable medium for
containing a portion of a test sample which is
located intermediate the first and second parts of
the housing and in substance-transferable contact
with the reaction medium when the first and second
15 parts of the housing are in a closed relationship in
a first closed position of the housing. The
reaction medium and medium containing an
analyte-indicating agent (or signal-producing agent)
are so positioned and arranged within their separate
20 housing parts that when the parts of the housing are
in a closed relationship in a second position, the
reaction medium and means for transferring the
analyte-indicating agent are in substance
transferable contact with one another.

25 In using this device, a test sample drawn from
a test subject is placed on a removable medium for
containing and transferring a portion of a test
sample which is placed in the two-part housing. The
housing is then closed to a first position such that
30 the removable test sample containing medium is in
transferable contact with and applies pressure to
the reaction medium. The housing is then opened and
the removable medium for containing and transferring

a test sample is removed from the housing. The housing is closed a second time to a second closed position such that the means for transferring an analyte-indicating agent is in transferable contact with and applies pressure to the reaction medium.

In another embodiment of the invention, various segregated areas of the reaction medium include a specific member of an immunological pair (MIP) which bind to or capture a specific analyte, such as a drug or drug metabolite. Different segregated areas may contain MIPs that are specific for different designated analytes, thereby allowing determinations for more than one analyte to be carried out in a single assay.

The methods and devices according to the invention are particularly beneficial in that only a small amount of signal-producing agent is required, in contrast to the conventional flow through cassettes or devices; applying pressure while maintaining direct contact on the surface of the test device maintains a localized concentration of reagents, particularly the signal-producing agent, which appears to increase the speed and sensitivity of the reaction, e.g., test results are obtained in a matter of seconds versus a matter of minutes or longer; and, in some embodiments, labeling a secondary antibody (as opposed to labeling the primary antibody or the antigen) decreases the reaction time of the assay because the antigen's shape is not altered by being bonded to a label.

DEFINITIONS

1. LIGAND-RECEPTOR ASSAY: any technique

involving the detection of the complex formed between a ligand and a substance which binds to the ligand. Preferably, one member of the complex is an analyte. The preferred ligand-receptor assay is an immunoassay. Ligand-receptor assays may be used to determine the presence, absence, quantity, and concentration of ligands in biological fluids. Ligand-receptor assays may be competitive or non-competitive, homogeneous or heterogeneous, direct or indirect, or a combined detection technique, e.g., binding an antibody to a small chemical moiety such as biotin or dinitrophenol (DNP).

It is preferred that during the assay process, substantially all of at least one predetermined ligand or ligand receptor remains in a predetermined position. Any technique for immobilizing a ligand or ligand receptor is included in the scope of the present invention. In a preferred embodiment, a ligand or ligand receptor is bound or immobilized on or in a solid phase. Typical immobilization mechanisms include, but are not limited to, covalent binding, noncovalent binding, chemical coupling, physical entrapment, and adsorption.

2. LIGAND: refers to the analyte itself, or a substance which can be used to infer the presence of an analyte in a test sample. A ligand-receptor refers to the substances for which there is a specific binding partner, e.g., the ligand. As used herein, ligand and ligand-receptors may be members of an immunological pair (MIP). Ligands and ligand-receptors may include haptens, hormones, antigens, antibodies (including anti-antibodies and

antibody fragments such as Fab or Fc), DNA, RNA, oligonucleotides, nucleic acids, and complexes or metabolites of any of the above. The ligand or the ligand-receptor may be labeled or unlabeled.

5 3. ANALYTE: the substance to be assayed, and, depending on the specific assay used, may be a ligand or a ligand-receptor. Exemplary analytes include but are not limited to drugs, proteins, haptens, hormones, metabolites of the
10 aforementioned, and other molecules, alone or in combination with a protein. A hapten is a small molecular weight material, such as some drugs or drug metabolites, which typically first attach to a protein in order to be antigenic or immunogenic.
15 For example, some drugs may be found in the body in both free and bound forms, which in turn provides the ability to distinguish between chronic drug abusers and acute drug abusers. In the case of the former, in many instances an antibody is formed in
20 the body against the bound form which can also be detected as an analyte by the present invention.

 4. BIOLOGICAL FLUID: any body fluid which may be tested to determine the presence or absence of an analyte, including, but not limited to blood
25 or a blood component, saliva, urine.

 5. Signal-producing AGENT: any agent used in a ligand-receptor assay which can be used to produce a detectable signal. The preferred signal-producing agents provide an easily visible signal, and more
30 preferably a color.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a plan view of an assay device

according to the present invention.

Figure 2 is a plan view of an assay device according to the present invention, illustrating the results of conducting a test for the presence or
5 absence of an antigen and the formation of a pattern suitable for identifying a test subject.

Figure 3 is an enlarged schematic cross sectional view of an assay device according to the present invention, showing a reaction zone in a
10 reaction medium.

Figure 4 is an enlarged schematic cross sectional view of an assay device according to the present invention, showing a control zone in a reaction medium.

15 Figure 5 is an enlarged schematic cross sectional view of an assay device according to the present invention, illustrating a positive test result.

Figure 6 is an enlarged schematic cross sectional view of an assay device according to the present invention, illustrating a negative test
20 result.

Figure 7 is an abstract representation of a cross section of an assay device according to the present invention, illustrating contacting the assay
25 device method according to the invention.

Figure 8 is an abstract representation of a cross section of an immunoassay device according to the present invention, illustrating the formation of
30 a fingerprint in a control zone of the reaction medium.

Figure 9 is an abstract representation of a cross section of an another embodiment of an

immunoassay device according to the present invention, illustrating the transfer of a test mixture from a swab to a finger of the person providing the test sample.

5 Figure 10 is an abstract representation of a cross section of the immunoassay device of Figure 9, illustrating the application of the test mixture to the reaction medium by the finger of the person providing the test sample.

10 Figure 11 is an open, exploded perspective view of one embodiment of a device according to the present invention.

15 Figure 12 is a perspective view of another embodiment of a device according to the present invention.

 Figure 13 is a perspective view of still another embodiment of a device according to the present invention.

DETAILED DESCRIPTION OF THE INVENTION

20 A ligand receptor assay device according to the present invention includes a reaction medium having at least one reaction zone and, preferably, at least one control zone which is capable of establishing the identity of a test subject. In a preferred
25 embodiment of the invention, the reaction zone includes at least one member of a ligand, ligand-receptor pair (also referred to herein as a "ligand/receptor pair"), this member being capable of establishing the presence or absence of a
30 substance in a test sample. In another preferred embodiment of the invention, the identity of the test subject is established by incorporating a

ligand in at least one control zone to provide for a test subject identifying pattern.

A method in accordance with the invention includes conducting, on or in a portion of a reaction medium, an immunological assay for the presence or absence of an analyte in a test sample, and separately establishing the identity of the test subject who provided the test sample on a portion of the reaction medium.

One embodiment of the invention is directed to an assay device, preferably an immunoassay device, comprising a porous support and a reaction medium covering a portion of the porous support. The reaction medium includes at least one reaction zone and, preferably, at least one control zone, wherein the control zone is capable of providing a pattern suitable for identifying a test subject. In a more preferred embodiment, the control zone is capable of providing a pattern comprising a fingerprint or surface reproduction of another part of a test subject's body which can be used for identification. The reaction zone includes a means for conducting a ligand/anti-ligand or ligand/ligand-receptor assay, preferably a means for conducting an assay for the presence or absence of an analyte. In another preferred embodiment, the means for conducting an assay includes a member of an immunological pair.

Another embodiment of the invention is directed to an assay device comprising a reaction medium having at least one reaction zone and, preferably, at least one control zone including a member of an immunological pair, wherein the control zone is capable of establishing the identity of a test

subject. The control zone is preferably capable of providing a pattern comprising a fingerprint or surface reproduction of another part of the test subject's body which can be used for identification.

5 According to another aspect, the present invention includes a support, preferably a porous support; a reaction medium, which is preferably a porous medium, mounted on the support, wherein the reaction medium has at least one reaction zone and
10 preferably at least one control zone, and wherein at least a portion of the control zone is capable of establishing the identity of the test subject. A preferred embodiment includes a member of an immunological pair positioned in or on the reaction
15 medium. In another preferred embodiment, at least a portion of a control zone includes one member of an immunological pair. In some embodiments of the invention, the reaction medium may be unsupported.

The present invention is also directed to a
20 method for processing a sample comprising contacting a reaction zone of an assay device with a test sample; determining the presence or absence of at least one analyte in the test sample; and forming a test subject-identifying pattern in a control zone
25 of the assay device.

The present invention is also directed to a method for determining the presence or absence of an analyte in a sample including conducting an
immunoassay for the analyte in a reaction zone of an
30 immunoassay device, whereby the presence or absence of an analyte in a test sample is determined; and establishing the identity of the person providing the test sample in a control zone on the immunoassay

device.

A method according to the invention includes determining the presence or absence of an analyte in a sample, preferably by using any ligand binding
5 based assay, and establishing the identity of the person who provided the sample, preferably by using any immunologically based protocol. Referring to Figure 1, the presence and/or absence of an analyte in a sample may be determined by applying a
10 biological fluid to at least one reaction zone 20, and allowing a predetermined ligand/receptor binding reaction to take place. In a preferred embodiment of the invention, completion of the reaction will result in the production of a signal. Preferably,
15 the signal is produced in the form of a visible color or the absence of color. For example, in Figure 2, the production of visible color hatched areas 32-34 indicates the absence of the analyte in the test sample, and the absence of color areas
20 35-37 indicates the presence of the analyte in the test sample. It is intended that the invention should not be limited by the specific binding assay employed, the method of producing the signal, the type of signal produced, and the particular location
25 of positive and/or negative results.

In accordance with the invention, the production of an image or pattern which identifies the person providing the test sample may be accomplished by applying an identifying body part, such as a
30 person's finger or foot, to a control zone, and allowing a predetermined ligand/receptor reaction to take place, thereby producing the image of a fingerprint or footprint. In a preferred embodiment

of the invention, the tip of a person's finger, coated with a signal-producing agent, preferably a color forming agent, is applied to the surface of a control zone having a MIP which will bond to the signal-producing agent coated on the person's finger. In a preferred embodiment of the invention, completion of the immunological reaction will result in the production of a visible color image which corresponds to the person's fingerprint. Figure 2 shows an exemplary visible fingerprint.

In an embodiment of the invention, particularly for pediatric use, the image or pattern which may be formed may be of the foot or a portion of the foot. In this embodiment, the foot or a portion of the foot, previously coated with a labeling reagent, may be applied to an apparatus according to the invention.

Exemplary embodiments of the invention will now be described by reference to the figures.

An immunoassay device 10 according to the invention includes a reaction medium 14, preferably mounted on a support or base member 12. Base member 12 is preferably a porous support. In a preferred embodiment of the invention, the reaction medium 14 includes at least one reaction zone 20 and at least one control zone 16. In the exemplary embodiments shown in Figures 1 and 2, the immunoassay device 10 includes six reaction zones 20 (numbered 32-37) and two control zones 16 and 18.

In a detection apparatus 10 according to the invention, at least one of the reaction zones 20 include means for conducting an immunological assay, preferably to establish the presence and/or absence

of an analyte in a test sample. In the exemplary embodiment shown in Figure 3, reaction zones 20 include at least one carrier, such as particles 22, embedded in reaction medium 14. As shown, the
5 particles 22 are coated with a member of an immunological pair 21.

In a detection apparatus 10 according to the invention, at least one of the control zones 16 include means for establishing the identity of a
10 test subject, preferably including a member of an immunological pair. In the exemplary embodiment shown in Figure 4, control zone 16 includes a carrier, such as particles 24, embedded in the reaction medium 14. As shown, the particles 24 are
15 coated with a member of an immunological pair 25. In a preferred embodiment of the invention, the MIP 25 is capable of bonding to a signal-producing agent, or to a MIP-signal-producing agent conjugate.

An exemplary method according to the invention
20 may be described by reference to Figures 5-8. In Figure 5, the presence of an analyte 31 in a test sample to which a MIP 30 has been added can be determined. The biological fluid is applied to the surface of reaction zone 20 having particles 22
25 coated with a member of an immunological pair 21. If analyte 31 is present in the biological fluid, it will bond to or be captured by MIP 30, with which it is mixed and incubated and which is specific for analyte 31. The resulting immunological pair
30 will not bond to or be captured by MIP 21, and will pass through the reaction medium 14. According to this embodiment of the invention and in this situation, no color forming agent will bond to the

coated particles 22.

In Figure 6, the absence of an analyte in a test sample to which a MIP 30 has been added can be determined. The biological fluid is applied to the surface of reaction zone 20 having particles 22 coated with a member of an immunological pair 21. In the absence of analyte in the biological fluid, MIP 30 will bind or be captured by MIP 21. According to this embodiment of the invention and in this situation, a color forming agent applied to the surface of the reaction medium will bond to MIPs 30 on the coated particles. In an alternative embodiment of the invention, the MIP 30 may include the color forming agent.

Figures 7 and 8 illustrate an exemplary method of producing the image of a fingerprint on the surface of a control zone. The surface 60 of a person's finger includes ridges 61 and valleys 62. When a person's finger is coated with a color forming reagent 102, a greater portion of the color forming reagent will tend to collect in the valleys 62. When the finger is applied to the surface of the reaction medium 14, it is believed that zones of higher concentration color forming reagent localize in the area of the valleys. The coated particles in the surface of the control zone will typically bond to a higher proportion of the color forming reagent, which will in turn produce a more intense visible color corresponding to the valley of the person's fingerprint.

In a preferred embodiment of the invention, determining the presence/absence of a substance in a biological fluid and producing the image of a

fingerprint are typically accomplished in only a few minutes.

Both the rapidity and higher sensitivity of the test devices and method of the present invention are attributable to characteristics of both the device and manner of applying reagents and/or test sample to the membrane surface and, to some extent, the materials from which the membrane is formed. Thus, as described below, the manner of application and materials are such that reagents in solution are maintained in contact with the membrane surface for a longer period of time to promote a greater degree of reaction between reagents in the membrane surface and reagents in solution. This leads to higher sensitivity and requires smaller concentrations of reagents.

Individual aspects of the invention will now be described in greater detail.

LIGAND-RECEPTOR ASSAY

In accordance with the invention, the detection device may be used with any assay technique for detecting an analyte of interest. In all of the assays used in accordance with the present invention, an analyte is bound to at least one ligand or ligand-receptor. Exemplary binding assays include but are not limited to labeled ligand assays, including competitive binding assays in which a solid phase is coupled to the binding reagent, such as simultaneous or sequential incubation with one or more separation steps, and displacement assays; and labeled binding reagent

assays, including noncompetitive and competitive binding assays, including assays in which the solid phase is the binding reagent or the ligand, and sandwich assays, including precipitation,
5 radioimmunoassay, or enzyme-linked immunosorbent assay. It is intended that the invention is not to be limited by the type of immunoassay employed or the specific protocol used in performing the assay. Exemplary immunoassays techniques are shown in the
10 Examples.

REACTION MEDIUM

In accordance with the invention, the reaction medium may be any medium suitable for performing a ligand-receptor assay. As noted above, the reaction
15 medium includes at least one reaction zone to indicate the presence or absence of an analyte and, preferably, at least one control zone to indicate the identity of the test subject. Typically, ligand-receptor assays may be conducted using a
20 reaction medium in the form of beads, tubes, plates, paddles, a porous or non-porous matrix, or porous or non-porous membranes or filters.

The reaction medium can include a depth medium wherein the surface or an area near the surface
25 comprises at least one reaction zone. The reaction medium may also include a layered structure having a reaction zone as a surface layer. In accordance with the invention, the reaction and control zones are positioned in the reaction medium so that a
30 signal-producing agent, if present, can be readily visualized.

In accordance with the invention, the preferred

reaction medium is porous, and even more preferred, a porous membrane. It is intended that the invention should not be limited by the type or construction of the reaction medium. One skilled in the art will readily recognize that the choice of reaction medium depends on the desired physical properties, including, but not limited to properties such as sensitivity, binding capacity, stability, the type of molecules or MIPs which can be bound, and compatibility with a particular assay protocol.

Exemplary materials for forming the reaction medium include, but are not limited to latex, nitrocellulose, nylon, cellulose, and nylon.

A preferred reaction medium is the commercially available Gelman Supor membrane. Supor membrane is a latex low protein binding polysulfone membrane with a hydrophilic surface. One skilled in the art will recognize that this membrane may be desirable because it provides superior flow rate and particle retention; a smooth surface; brilliant whiteness and opaqueness to enhance signal contrast in diagnostic tests; low extractables to reduce sample contamination and background interference; and uniform porosity to ensure final product consistency. Also, the use of this membrane obviates the need for an external wetting agent, which may be desirable for controlling the introduction of unwanted extractables.

Another preferred membrane, IAM, Immobilon Affinity Membrane (commercially available from Millipore Corporation; see, for example, U.S. Patent 4,407,943), the hydrophobic base polymer, polyvinylidenedifluoride (PVDF), binds proteins.

IAM also permits a high degree of control over the extent of protein binding and the user can reproducibly immobilize nanogram to microgram quantities of protein on the surface to suit various assay requirements. Binding a ligand to IAM and its use in immunoassays is known in the art.

The use of membranes, as the solid support in a reaction medium of the present invention may be desirable for the added convenience of a solid phase, the typically high protein binding capacity, and the membrane's flow-through characteristics. Nitrocellulose is one of the most commonly used membranes due to its high affinity for proteins nucleic acids, other cellular antigens, and cellular macromolecules, and may be desirable if the desired MIP binding is ionic or hydrophobic. Nitrocellulose also provides an excellent matrix for blotting proteins and nucleic acids. The nitrocellulose may be cut into whatever shape is required and has the useful characteristic that the amount of protein in a fingerprint will be clearly visible.

Included within the scope of the present invention is changing or incorporating different surface properties on the membrane in order to achieve a desired result, e.g., the surface properties of a membrane designed for a competitive binding assay for a hormone may be different than an immunometric assay for a therapeutic drug. For example, it has been shown that treating the surface of a hydrophilized PVDF membrane with ethanolamine reduces the non-specific binding of the membrane surface. Selection of a particular surface treatment agent or surface property may be based on

the desired chemical characteristic to be imparted to the surface; the inability or reduced capability of denaturing or impairing the functionality of a bioactive agent on or in the reaction zone; the
5 desire to effect a certain orientation of an immobilized bioactive molecule; the desire to promote long-term stability of an immobilized bioactive molecule; the inclusion of a desired nucleophilic substituent; and the availability and
10 cost of treatment agents. The use of other surface treatment agents, including bifunctional or multifunctional reagents, to affect the surface properties of the membrane are included within the present invention.

15 One skilled in the art will readily recognize the benefit derived by including a reaction medium which is porous. Positive test samples will readily pass through the reaction medium, obviating the need to wash the surface of the reaction medium prior to
20 adding a color-forming agent. Although it is less desirable, it is intended that the invention include the use of a non-porous medium. A preferred support material is a somewhat rigid porous plastic material known as Porex.

25 As noted above, the reaction medium may be supported or unsupported. In the preferred embodiment, the reaction medium is supported, and even more preferred, the support is porous. An exemplary support includes, but is not limited to
30 polystyrene.

In another embodiment of the invention, the support may be impermeable to the fluids and reagents used in conducting the testing; in this

embodiment, the reaction medium is preferably "deep" enough to have a reaction zone at or near the surface of the medium.

In preferred embodiments of the present invention
5 in which the device is totally or substantially self-contained, that is, needing no additional reagents or with at most a few drops of a buffer solution or water to rinse the reaction medium, and needing only a minimum level of skill to perform an
10 accurate determination, another type of material is preferred to support the reaction medium. For such applications, a compressible wicking or sponge-like material is preferred. Thus, by applying pressure to the outer, substantially planer surface of the
15 reaction medium, which is adjacent, in contact with or bonded to a substantially planer surface of the compressible support material, causes compression of or an indentation in the surface of the support material adjacent the reaction medium, such that the
20 support surface is caused to be below its uncompressed planar surface. In the presence of a liquid, after pressure is released from the compressible support material and as the compressible material expands to its uncompressed
25 state, the liquid is drawn into the support material. Thus, if a liquid sample, reagent solution, buffer solution or water is applied to the reaction medium and sufficient pressure is applied to cause a deformation in the surface of the support
30 material, liquid will be pulled into the support material as it expands to its original state. The pressure applied is a function of the compressibility of the material. A preferred

compressible support material is a compressible, flexible reticulated urethane foam, preferably a polyester or polyether urethane foam having a pore rating of about 75 to 100 ppi, preferably about 80
5 to 90 ppi. Such materials are available from E.M. Murray Co. Inc.

CONTROL ZONE

In a preferred embodiment of the invention, the reaction medium includes a control zone for
10 establishing a reference point in determining the presence or absence of an analyte in a test sample (reference control zone), and most importantly a control zone for establishing the identity of the test subject who provided the test sample (identity
15 control zone).

In accordance with the invention, a control zone for establishing the identity of the test subject (identity control zone) who provided the test sample may include any signaling mechanism which
20 incorporates a MIP. For example, the control zone may include a member of an immunological pair capable of bonding to a signal-producing agent, which when applied by contacting the test subject's finger to the surface of the identity control zone,
25 an image of the fingerprint is produced on the test device.

As used herein, "capable of establishing the identity of a test subject" refers to the capability of specifically identifying the subject who provided
30 the sample being tested. In a preferred embodiment of the invention, the control zone includes a member of an immunological pair which, when placed in

contact with the test subject's finger which has been previously coated with a signal-producing agent, is capable of producing a permanent image of a fingerprint, preferably a fingerprint image easily visible. In another embodiment, the invention includes a control zone having a member of an immunological pair or nucleotide sequences which are capable of bonding to nucleotide sequences which can be used to identify the test subject.

In accordance with the invention, the control zone for establishing a reference point in determining the presence or absence of an analyte in a test sample includes any means for providing a control. Several examples for providing a control are shown in the examples. It is intended that the invention not be construed as being limited by the means or mechanism for establishing the standard of comparison in judging the test results.

REACTION ZONE

The reaction zone refers to the locus in which the presence or absence of an analyte can be determined. For example, the reaction zone is preferably on or near the surface of the reaction medium, or may be below the surface. In a preferred embodiment of the invention, the reaction zone includes at least one ligand or ligand-receptor bound to a particle or bead in a matrix or filter, suitable for capturing an analyte of interest. In another embodiment of the invention, at least one of the reaction zones may be a control for establishing a reference point, e.g., a comparative control to show the absence of color.

Signal-producing AGENT

A signal-producing agent refers to any agent or marker which produces a detectable signal or which permits the detection of a ligand or

5 ligand-receptor. Preferred signal-producing agents are those which permit detection of the analyte without instruments, preferably by visual means. Exemplary signal-producing agents include, but are not limited to color forming agents, such as an

10 enzyme, polymer containing dyes, chemiluminescent agents, fluorescent agents, radioisotopes or ferromagnetic particles. The color forming agent may be a colored particle, a colored molecule or some species, such as an enzyme, which is capable of

15 triggering a sequence of events leading to the formation of a colored marker. The colored molecule may be a fluorescent dye, such as fluorescein or rhodamine; a chemiluminescent compound; a bioluminescent compound; or a compound that may be

20 detected by the absorption of electromagnetic radiation (and possible reemission of radiation at another wavelength), including ultraviolet radiation, visible radiation and infrared radiation. The colored molecule may be directly or indirectly

25 conjugated to a ligand or ligand-receptor. Alternatively, the colored molecule may be incorporated in a particle, particularly a microsome.

Enzymes, useful as color forming agents, include

30 alkaline phosphatase, horseradish peroxidase or B-galactosidase. Such enzymes are often used in conjunction with a chromogenic substrate. A list of some exemplary chromogenic substrates which may be

used with enzyme color forming agents are given below in Example 11.

In a preferred embodiment of the invention the color forming agent may be an electron dense
5 particle, such as colloidal gold, silver coated colloidal gold or ferritin.

In another preferred embodiment of the invention, the color forming agent is capable of bonding to a MIP.

10 The present invention includes a signal-producing agent, which may be a color forming agent. Preferably, the signal-producing agent comprises a labeled MIP, for detecting the adsorption of an
15 analyte or an analyte conjugate in or on the reaction medium. The labeled MIP may comprise any one of a number of appropriate labels, such as an enzyme, a fluorescent compound, a bioluminescent compound, a ferromagnetic atom or a colored particle or microsome. In a preferred embodiment, the label
20 used in the present invention is colloidal gold. Gold is biologically inert, has very good charge distribution and is widely available in many useful forms. Its detection can be enhanced using several silver deposition methods, which permit development
25 to be monitored by the naked eye. Colloidal gold particles conjugated with a wide range of anti-immunoglobulin antibodies, protein A or streptavidin are available commercially.

While a colloidal gold substrate is preferred
30 over other dyed particles or microsomes, a chromogenic substrate provides an alternate sensitive detection method for the enzyme conjugate.

Several embodiments of the present invention

relate to compact assay devices and kits which contain all or most of the reagents and materials necessary for qualitative and/or quantitative determinations of one or more analytes. A preferred embodiment of such a device is illustrated in Figure 11 which shows a self-contained assay device 110. The device includes a housing 111 which in a preferred embodiment includes a first part 111a and a second part 111b. Located within the first part of the housing 111a, mounted on the upper surface of a raised plateau or platform portion of a base member 112 is a reaction medium 114, of the type described above. The base member 112 preferably includes a porous support adjacent, in contact with or bonded to the reaction medium 114. In a preferred embodiment of the invention, the reaction medium 114 includes one or more reaction zones 120, each of which includes a specific reagent, such as a member of a ligand/receptor pair (which may be a member of an immunological pair, for example an antigen or an antibody). The reaction medium in this preferred embodiment also includes at least one control zone 116. As in the embodiments discussed above, each of the reaction zones includes a reagent suitable for determining the qualitative presence or the quantitative range of an analyte. In a preferred embodiment of the invention, a control zone is suitable for identifying the donor of a test sample. Likewise, the control zone 116 also includes a suitable test reagent for identifying the test subject which is preferably a member of a ligand/receptor pair, such as a member of an immunological pair.

The reagents present in each of the reaction zones may be applied manually or, preferably, using an automated process, such as using an ink jet sprayer of the type used in ink jet printing systems, to apply the reagent in individual and segregated reaction zones.

In the second part of the housing 111b, is located a medium 126 on which is spread at least one signal-producing agent, or analyte-indicating agent, which is preferably a color forming agent. The signal-producing agent-containing medium 126 is formed from a material which is of a lower porosity than either the reaction medium or the medium for containing a test sample (discussed below). Preferable for use as the medium on which the signal-producing agent is spread is a closed cell polyolefin material, such as a closed cell polyethylene or polypropylene. A preferred material is available as Minicell from Voltek Corporation. The signal-producing agent, as with the other embodiments of the present invention, preferably is a labelled member of a ligand/receptor pair, such as a labelled member of an immunological pair, such as a colloidal gold labelled substance. The medium 126, containing the signal-producing agent, is positioned in the second part of the housing 111b in such a way that when the two parts of housing are brought together in a specific closed position, contact is made between the medium 126 and the reaction medium 116 such that signal-producing material contained in the medium 126 is transferred to the reaction medium. This is preferably accomplished by mounting the medium 126 on a base

portion 128 which resembles an elevated plateau or platform, which in the preferred embodiment shown in Figure 11 has a shape similar to base 112. This arrangement, in which the medium containing the
5 signal-producing agent is spaced apart from the reaction medium 114 in a first or open position and transfers the signal-producing agent to the reaction medium 114 in a second or specific closed position, apparently achieves greater sensitivity and
10 increased speed due to the pressure applied when the signal-producing agent-containing medium 126 contacts the reagent medium 114. Depending upon the specific materials involved, a pressure in excess of atmospheric may be used, preferably slightly less
15 than about 2 to about 5 psi, for a period of about 5 to about 20 seconds is sufficient to adequately retain the signal-producing agent at or near the surface of the reaction medium after the media 114 and 126 are placed in contact with one another.
20 Although higher pressures can generally be used to transfer either the signal-producing agent, analyte or conjugate formed of the analyte with a ligand receptor of the analyte (as discussed below), the duration of time for exerting the pressure is
25 decreased, since pressure and time vary inversely.

Another component of the self-contained or "integrated" device illustrated in Figure 11 is a medium 185 for containing and transferring a portion of a test sample to the reaction medium 114. The
30 medium 185 used to hold both a test sample and a reagent capable of reacting with an analyte is a hydrophilic material, preferably in the form of a hydrophilic sponge and preferably is formed from a

fine pored polyester urethane sponge material. A preferred material is polyester urethane foam sponge material having a porosity of 100 ppi that is available from E.N. Murray Company Inc. The medium 185 serves several purposes. First, it is capable of absorbing a test sample taken from a donor or test subject. It is also capable of transferring the sample or a conjugate formed from an analyte present in the test sample and a receptor against the analyte to the reaction medium 114. The medium 185 is also capable of providing diffusion and reaction loci for both the substances present in the test sample, as well as a reagent present in the medium. This reagent is preferably a member of a ligand/receptor pair, such as a member of an immunological pair of the type discussed above and identified in Figures 5 and 6 as MIP 30. The reagent in the medium 185 is typically an antibody. This reagent is contained in the medium 185 preferably as a lyophilized antibody. In this embodiment, the sponge medium 185 is treated with a solution of reagent and the treated sponge is subjected to a lyophilization treatment in a lyophilization chamber, such as a Speed-Vac.

The medium for containing the test sample and the test sample and the reagent, preferably in lyophilized form, may be mounted at one end of a cylindrical body 140. In the embodiment illustrated in Figure 11, the open end 140a opposite to the closed end 140b of the cylinder on which the medium 185 is mounted, is larger in shape or diameter than the base portion 128 of the housing part 111b on which the signal-producing agent-containing medium

126 is mounted. Preferably the open end 140a of the cylinder 140 is commensurate in shape to the base portion 128 so that the cylinder may be easily accommodated in the housing part 111b over the base portion 128. The cylinder 140 is of such a shape and configuration that in an "open" position of the housing parts in which the media 114 and 126 are in spaced-apart relationship with one another, the medium 185 is also in spaced-apart relationship with both of the media 114 and 126 in the open or first position of the housing members 111a and 111b. In a second or specific "closed" position of the housing members 111a and 111b, while remaining within the housing 111 in and spaced relationship to the base 128 and signal-producing agent-containing medium 126, the medium 185 for containing the sample contacts and transfers substances present in the sample and/or a ligand/receptor pair formed between an analyte found in the sample and a conjugate thereof to the reaction medium 114. Thus, in the second or closed position, pressure is exerted by the end of the cylinder 140b and medium 185 on the reaction medium 114 in much the same manner as in the contact which occurs between the medium 126 containing the signal-producing agent and the reaction medium in a second closed position. The method of using this device is discussed below.

While Figure 11 illustrates an embodiment of the present invention which employs a two-part housing 111 having a first part 111a and a second part 111b, and a removable member 140 on which a sponge 185 for containing a test sample is attached, other aspects and alternative embodiments are contemplated by the

present invention. For example, the housing parts 111a and 111b are illustrated in Figure 11 as being joined to each other by a flexible hinge arrangement. When the housing is formed from a
5 suitable plastic, which is preferably substantially inert to any of the reagents or materials used in other components of the device or in solutions applied to the media of the device, the hinge may be formed from a thinner portion of the same plastic
10 material. Thus, the housing may be formed as a single unit in one pressing operation. Alternatively, another material may be used as the hinge material and the hinge attached to the housing parts by suitable means.

15 In another embodiment, the housing parts 111a and 111b need not be joined to one another at all. Thus, to close the device, the separate parts may be constructed to snap into place with a detent or some similar arrangement or may engage one another with a
20 threaded arrangement or a bayonet engaging arrangement.

In yet another alternate embodiment, the medium 185 for containing the sample and containing a test reagent, rather than being mounted on the removable
25 cylindrical member 140 of the type shown in Figure 11, may be located in a third housing part. When used with housing parts which are permanently joined to one another, such as that described immediately above, the housing part containing the sponge for
30 containing a test sample may be placed intermediate the separate housing parts containing the reaction medium and the signal-producing agent-containing medium (equivalent to housing parts 111a and 111b).

In this instance, however, the intermediate housing part containing the sponge 185 may engage one or both of the other housing parts and contact the reaction medium in the first housing part when in place and forming part of the housing. It may also be removed when it is desired to contact the reaction medium 114 with the medium containing a signal-producing agent 126.

In yet another embodiment, the housing part containing the hydrophilic sponge for containing a test sample may be joined to the first housing part (equivalent to housing part 111a as shown in Figure 11) by means of a hinge on the edge portion of the first housing part in opposing relationship to the edge of the first housing part to which the second housing part is joined. Thus, when it is desired to place the sponge containing a test sample in contact with the reaction medium, the third housing part containing the sponge may be rotated to a substance-transferring position in contact with the reaction medium and thereafter rotated to a spaced apart position with respect to the reaction medium and the second part of the housing containing the signal-producing agent medium 126 may be rotated into transferable contact with the reaction medium.

The above embodiments of the present invention which include three media (a reaction medium, a medium for containing a test sample, and a medium containing a signal-producing agent) are substantially self-contained. Thus, it is unnecessary to add additional reagents in order to perform the assay. At most, a buffer solution, such as a phosphate buffered solution, or water may be

used in a final step to rinse the reaction medium to remove residual reagents from the surface of the reagent medium. Alternatively, when a compressible porous medium is used as a support for the reaction medium, it may be unnecessary to use any supplemental materials.

The present invention also contemplates compact assay devices using only two media, rather than the three-media devices discussed above. Thus, in the embodiment illustrated in Figure 12, while still retaining the reaction medium 114 mounted on base 112 in a first part of the housing, the sample-containing and transferring sponge 185 may be mounted directly on the base 128 instead of the signal-producing agent-containing medium 126 being located thereat. In such an embodiment, the cylinder 140 on which the medium 185 is mounted in the "3-media" embodiment illustrated in Figure 11 may be dispensed with. In such an embodiment, the wash bottle containing a buffer, detergent or water may be used in the same manner as described above. However, in this embodiment, the signal-producing agent may be dispensed from a reagent bottle directly on the finger or other identifying body part of the test subject and the finger or other body part may be placed in contact with the reaction medium after the test sample has been placed on the sponge 185 mounted in base 128, incubation has taken place and the housing has been closed such that there is substance transferable contact with suitable pressure between the surface of sponge 185 and the reaction medium 114.

Another two-media device is illustrated in Figure

13. Like the embodiment discussed above and shown
in Figure 11, this includes the reaction medium 114
mounted on base 112 in a first part of the housing
and the signal-producing agent-containing medium 126
5 in the second part of the housing. This corresponds
substantially to the embodiment shown in Figure 11
without the cylinder 140 and medium 185 for
containing a test sample. This embodiment, like the
3-media device illustrated in Figure 11 does not
10 require the presence of the test subject, and,
accordingly, will not produce a fingerprint or other
identifying mark of the test subject when the test
subject is not present. Alternatively, however, a
fingerprint can be produced when the test subject is
15 available at the time the test is performed. In
this embodiment, the sample is mixed with a test
reagent supplied with the device, most suitably in a
reagent bottle, as part of a kit. The reagent may
include one or more substances capable of indicating
20 the presence of one or more analytes and is
preferably a member of a ligand/receptor pair, most
preferably a member of an immunological pair capable
of forming a conjugate with a specific analyte. The
mixture of test sample and reagent is applied
25 dropwise to the surface of the reaction medium,
sufficient time for incubation allowed and the
housing is closed for a short period of time so that
there is contact between the signal-producing agent-
containing medium 126 and the reaction medium.
30 sufficient to transfer the signal-producing agent to
the surface of the reaction medium. If the test
subject is present at the time of testing, rather
than closing the device to make contact between the

media 114 and 126, the test subject may roll a finger on the surface of the signal-producing agent medium 126 and then press the finger on the reaction medium.

5 Another embodiment of the present invention is directed to a device and method for distinguishing between an acute and a chronic drug abuser. Thus, in the body of one who uses a drug, not only are molecules of the drug or a metabolite found, but the
10 drug or such metabolite binds to a carrier molecule. Continued use of a drug stimulates production of an antibody against the carrier bound drug. Thus, in the body of a human who chronically uses a drug may be found not only a residue of the drug or
15 metabolite thereof, but also the presence of a human antibody to the drug. In an embodiment of the device according to the present invention, a reaction zone in the reaction medium may be spotted with some of the drug of which the test subject is
20 suspected of using. This will then form a conjugate with the human antibody to the drug if present as an analyte in a test sample. As a signal-producing agent for indicating the presence of the drug antibody may be used the corresponding goat or mouse
25 antihuman labelled antibody.

METHOD OF USE:

An assay method in accordance with the invention includes conducting a ligand-receptor assay for an analyte in a reaction zone of an assay device,
30 whereby the presence or absence of an analyte in the test sample is determined, and establishing the identity of the person providing the test sample in

a control zone on the assay device. In a preferred embodiment of the invention, establishing the identity of the person providing the test sample may be accomplished using ligand-receptor assay methods, e.g., binding a MIP to a signal-producing agent.

As noted above, the biological fluid sample to be tested for the presence or absence of at least one analyte is applied to the surface of the immunoassay device according to the invention. Any means of applying the fluid sample to the detection apparatus may be used. For example, in one embodiment of the invention, the biological fluid, e.g., urine, may be combined with a MIP according to standard immunoassay procedures, drawn into a pipette, and deposited on the surface of the test device.

In another example, the biological fluid, e.g., blood, may be collected in a pipette and deposited on the surface of the test device, or may be smeared on a solid applicator, such as a fingertip. In one preferred embodiment, blood is smeared on the test subject's fingertip, which is then pressed in contact with the surface of the detection device. In another preferred embodiment, the biological fluid is combined with a MIP before application to the reaction medium. In still another preferred embodiment, the sample is deposited on the surface of a medium for containing a test sample. Other exemplary methods of applying the test sample to the detection device are shown in the Examples.

A preferred embodiment of the invention includes applying one or more reagents or a sample to a surface of the test device using pressure. For example, it has been found that smearing a

signal-producing agent on the test subject's fingertip and applying the fingertip to the test device significantly reduces the amount of signal-producing agent required, and significantly
5 increases the sensitivity and speed at which the assay proceeds. For example, as little as 10 microliters and preferably about 12 microliters of colloidal gold can be smeared on the test subject's fingertip in order to perform the assay. In
10 contrast, as much as 100 microliters or more of the signal-producing agent is typically required for conventional flow-through cassette assay devices. In a like manner, another preferred embodiment includes the use of pressure in the application of
15 sample to the surface of the reaction medium.

While not intending to be limited to a particular theory of operation, it is believed that the pressure and/or surface tension which occurs when the test subject's finger is placed on the assay
20 device results in localizing an increased concentration of ligand and ligand receptor, increasing the amount of time that the reagents are retained in a localized area of the assay device, and may involve migration of the signal-producing
25 agent from the ridges of the fingerprint to the valleys, to produce a clear fingerprint pattern on the surface of the test device.

The higher sensitivity and reduced amounts of reagents provided by the present invention seems to
30 result from a more complete reaction taking place at the surface of the membrane where immobilized reagent is found. The more complete reaction results apparently from the reagents applied to the

surface of the membrane in liquid form being maintained thereat for a longer period of time. Thus, as is typical with most known test devices, liquid samples and/or reagents are applied in drop
5 form to the surface of a membrane which contains either reagent or test sample and are quickly drawn through the membrane, either because of an absorbent placed in contact with the membrane on its downstream surface or because of the nature of the
10 membrane material. In the present invention, however, the liquid containing the reagent and/or sample is applied to the surface of the reaction medium and is maintained thereat with the application of pressure greater than atmospheric
15 pressure as a thin film which shows little tendency to pass completely through the membrane. Thus, using the finger as an applicator or another applicator means, such as an applicator means included as part of the device, reagent-containing
20 liquid on the applicator is transferred to and maintained at the upstream surface of the reaction medium under pressure. Thus, while not wishing to be held to any theory, it appears that primarily the surface tension of the liquid, in combination with
25 other factors such as the capillary action of the applicator and possibly the porosity of the medium on which the liquid is placed in combination with the pressure causes a thin film of liquid to be maintained at the upstream surface rather than
30 passing through the medium as long as the pressure is maintained on the outer surface of the medium. The contribution of the surface tension in maintaining a thin film at the surface of the

reaction medium appears to increase the availability of the substance(s) present in the liquid to reagents in the medium. Thus, using an applicator, such as the finger, with a pressure greater than
5 atmospheric pressure, typically as little as about 1 to about 5 psi and up to a pressure that a test subject would normally and comfortably apply (such as in pressing the finger in making an "official fingerprint"), prevents the liquid from passing
10 through the membrane until the pressure is removed. Application of pressure would normally be for a period of time of about 5 to 20 seconds. Maintaining the reagent at the surface of the medium or membrane for this period of time is sufficient to
15 assure a greater degree of reaction of the reagents located at or immediately below the upstream surface of the medium. While the nature of the liquid, such as its surface tension, as well as the materials from which the membrane and applicator are formed,
20 have some bearing on this phenomenon, it appears that a variety of materials may be used and the phenomenon of maintaining the thin film of liquid at the surface of the membrane still occurs. Furthermore, and also not wishing to be held to a
25 particular theory, it appears that the increased sensitivity and particularly the rapidity of the assay determinations of the devices and methods of the present invention may be attributed, at least in part to the use of pressure at the time liquid
30 solutions are applied to the surface of the reaction medium. It is believed that the pressure applied, which is greater than atmospheric pressure and is typically as little as 1 psi and preferably in the

range of about 2 to 15 psi, increases the kinetics or the speed of the particular reaction taking place.

The test device preferably includes one or more
5 MIPs immobilized in a corresponding number of reaction zones. For example, if the analyte of interest is an antibody, the test device may include an antigen bound to microspheres or particles embedded in the reaction zone. One skilled in the
10 art will recognize that a desired amount of antigen can be immobilized in the reaction zone, and that a threshold concentration of bound MIP may be used to detect a predetermined amount of analyte. In one embodiment of the invention, the detection device
15 includes several reaction zones, each reaction zone for a different analyte, and each reaction zone includes a predetermined threshold concentration of bound MIP to detect a predetermined amount of analyte. As used herein, threshold amount or
20 concentration refers to the lower limit of a concentration range for an analyte. In another embodiment of the invention the detection device includes several reaction zones, each reaction zone including a different threshold concentration for
25 the same analyte. In this embodiment, the threshold concentration provides a reference point for determining the upper limit of an analyte concentration range. As shown in the examples below, varying the threshold concentration in the
30 reaction zones may provide an exemplary method of quantifying the amount of analyte in a sample.

After the biological fluid being tested has been applied to the surface of the detection device, the

presence or absence of the analyte in the test sample can be determined, preferably by visually ascertaining a color (signal) or the absence of a color (signal). In a preferred embodiment of the invention, a positive response, i.e., indicating the presence of the analyte in the biological fluid, corresponds to the absence of color. In a preferred embodiment of the invention, a negative response, i.e., indicating the absence of the analyte in the biological fluid, corresponds to a visualized color. However, in one embodiment of the invention in which a determination is made as to whether the test subject is a chronic drug abuser in which a human antibody to a drug is being tested for and the reagent present in the reaction zone is the drug itself and the labelled signal-producing agent is, for example, a gold labelled goat or mouse antihuman antibody, a positive response, indicating the presence of the drug antibody, corresponds to the appearance of color.

In accordance with one aspect of the invention, when the biological fluid being tested is blood, it has been unexpectedly found that whole blood may be directly applied to the surface of the test device without the need for serum separation prior to the sample being applied to the test device. In accordance with this embodiment of the invention, the whole blood sample is applied to the surface of the test device, and then a detergent is applied over the blood sample. While it is known that detergents such as Tween may be used to remove red cells and the like from whole blood samples, it has been unexpectedly found that the application of a

clearing reagent, such as a detergent (preferably a detergent which includes a mixture of ionic, and nonionic detergents, and an alcohol), allows the application of whole blood to the surface of the assay device without the need to separate the cellular component from the serum prior to application to the device. The application of a detergent is preferable because the detergent disrupts blood cells. The ruptured cell debris then pass through the medium and are removed from the reaction zone, i.e., removed from interfering with immobilized surface antigens.

Once the presence or absence of the analyte has been determined, the detection device made in accordance with the present invention may be used to identify the test subject whose biological fluid was tested. In a preferred embodiment of the invention, determining the identity of the test subject includes performing a ligand-receptor assay which results in the development of a fingerprint of the individual. In accordance with a preferred embodiment of the invention, determining the identity of the test subject is independent of any reactions with the analyte in the fluid.

In accordance with this method of the invention, the person's fingertip is coated with a signal-producing agent conjugated to a MIP. When the coated fingertip is applied to the surface of a control zone, the signal-producing conjugate binds to a MIP embedded in the control zone of the test device. While the invention is not to be limited to a particular theory of operation, it is believed that the ridges and valleys of a person's fingertip

provide localized areas of concentrated signal-producing conjugate (corresponding to a valleys) and localized areas of low concentration signal-producing agent (corresponding to a ridge). When
5 applied to the surface of a control zone of a device according to the invention, a clearly identifiable image of a fingerprint is produced, whereby the identity of the person providing the test sample can be provided.

10 In accordance with the invention, the presence or absence of an analyte of interest in a sample may be determined by applying the sample to the surface of a reaction medium, the reaction medium including at least one reaction zone having a member of an
15 immunological pair therein; and applying a signal-producing agent which binds, directly or indirectly (e.g., a secondary antibody), to the MIP or to the analyte of interest.

The self-contained test device illustrated in
20 Figure 11 may be used as follows:

A test sample in liquid form, such as a bodily fluid (for example, saliva, urine or blood) is placed on the outer surface of the medium 185 for containing and transferring the test sample. Up to
25 about 20 drops of test sample may be applied to the medium 185. Since this device provides so much greater sensitivity and speed than do conventional test devices, as little as one drop of test sample may be employed preferably, however, about 3 to
30 about 10 drops of test sample are preferably used. With the cylinder 140 in place in the housing disposed with the open end thereof over the base 128 and signal-producing medium 126 in the first part of

the housing 111b, the test sample is applied to the outer surface of the medium 185 or the sample may be applied to the medium and the cylinder 140 thereafter placed in housing part 111b. Any analyte
5 which is present in the test sample undergoes reaction with a member of a ligand/receptor pair, preferably a member of an immunological pair, present in lyophilized form in the sponge 185, to form a conjugate. A period of about 1 to about 4
10 minutes, preferably about 2 minutes, is permitted for incubation (reaction) of the test sample and member of the ligand/receptor pair. With the cylinder 140 remaining in place in the second part 111b of the housing, the two parts of housing 111
15 are brought together to a first closed position in which the outer surface of the sponge 185 contacts the reaction medium 114 and exerts sufficient pressure to transfer substances present in the test sample, the member of the ligand/receptor pair and
20 any conjugate formed between the analyte and the member of the ligand/receptor pair. In this position, although being referred to as a "closed position", the housing may not be entirely closed, depending on the particular structure employed.
25 This closed position is maintained for a period of about 1 to about 4 minutes, preferably about 2 minutes. Thereafter, the housing is opened and the cylinder 140, containing the sponge 185 is removed from the housing device. The device is then closed
30 to a fully closed or second closed position in which the signal-producing agent-containing medium 126 contacts the reaction medium 116 with a sufficient pressure to transfer the signal-producing agent,

such as a gold-labelled member of a ligand/receptor pair to the reaction medium. The housing is maintained in this second or fully closed position for a period of at least about 5 to about 30

5 seconds, preferably about 15 seconds.

After about a minute the device will indicate the presence of any analytes being tested for, or an amount of analyte being present in excess of a predetermined concentration or amount, by showing a
10 blank space in the appropriate reaction zone. In most applications, a negative test is reflected by a coloration of the appropriate reaction zone when a specific analyte is absent or is present in amounts where concentration is below a predetermined value.

15 A device will not provide an indication of the identity of the test subject at this stage unless the test subject places a finger or other identifying portion of the body in contact with the control zone 116 of the reaction medium 114 after
20 opening the housing for the second time.

Alternatively, rather than applying the finger or other body part to the control zone of the reaction medium after contact has been made between the medium 126 and 114, this step of pressing the two
25 media together may be omitted and the finger or other body part of the test subject may contact under pressure the reaction medium after first being rolled over the signal-producing agent-containing medium 126 or applying the signal-producing agent
30 directly to the finger or other body part with an applicator or dispenser. While each of these methods of identifying the test subject by a permanent and individual characteristic of the test

subject is both rapid and sensitive (the total determination typically taking no more than about 3 to 5 minutes), the last described method in which the signal-producing agent is applied directly to the fingertip which is then applied to the reaction medium, is somewhat more effective in that smaller amounts of reagent and sample may be required and the speed of the process is somewhat more rapid. The method in which a finger or other identifying body part is not placed in contact with the reaction medium is least effective in terms of the amounts of material required and speed of the process while contact between the finger or other body part with the transfer medium is intermediate in effectiveness.

TEST KIT

In an embodiment of the invention, the assay device made in accordance with the invention may be incorporated into a kit. The kit may also include any of a number of reagents for performing the ligand/receptor assays. For example, the kit may include a test device, at least one agent for producing a signal (or an "ink" pad for transferring the signal-producing agent to the finger), and one or more washing, clearing or detergent reagents. The kit may also include one or more ligands or ligand receptors, such as lyophilized primary antibody and a secondary antibody, and one or more buffers. The ligand or ligand receptor may be a labelled material.

When the compact devices are supplied in kit form, they may be accompanied by a wash bottle

containing a phosphate buffer solution. Such wash
bottle and/or reagent bottle may be supplied most
effectively as a kit with the alternate two-media
embodiments of the device of the present invention
5 shown in Figures 12 and 13, discussed above. Thus,
rather than the device illustrated in Figure 11
containing a housing and the three media, 114, 126
and 185, a device including a housing and two of the
three media, and appropriate reagents contained
10 therein may be employed.

SPECIFIC EXAMPLES

EXAMPLE 1.

One embodiment of a immunoassay device of the present invention was prepared according to the following procedure. A reaction zone was constructed by embedding polystyrene particles, coated with human serum albumin-benzoylecgonine (HSA-BE), in one portion of a low protein binding polysulfone membrane (Gelman Supor membrane) mounted on a porous support. A control zone 16 was constructed by embedding polystyrene particles, coated with goat anti-mouse antibody, in a different portion of the membrane. A representative schematic cross section of the reaction and control zones is shown in Figures 3 and 4.

The assay device was prepared according to the following procedure. The reaction medium was first rinsed with a blocking buffer solution (Solution A) consisting of 2% polyvinyl alcohol (PVA), 1% glycine and 0.05% Tween-20 in a phosphate buffered saline solution. The central well of the immunoassay device (control zone 16) was spotted with a dilute solution of polystyrene latex coated with goat anti-mouse immunoglobulin G (adsorbed against human serum) in a phosphate buffered saline containing 4% sucrose, 1.0% BSA and 0.05% azide. The reaction zone was spotted with a dilute solution of polystyrene latex coated with human serum albumin-benzoylecgonine (HSA-BE) in 0.2 M sodium bicarbonate. The reaction medium was then inverted onto hydrophobic polyethylene and dried for one hour

in a drying room set between 80 and 100°F. After removal of the reaction medium from the hydrophobic polyethylene, the assay device was ready for use.

EXAMPLE 2.

5 A saliva sample was obtained from a person to be tested for the presence or absence of benzoylecgonine. A 200 microliter aliquot of a dilute solution of mouse anti-benzoylecgonine immunoglobulin G (mouse anti-BE IgG) in a phosphate
10 buffered saline, which contained 0.1% bovine serum albumin (BSA) and 0.05% Tween-20, was added to a 200 microliter aliquot of the saliva sample. The resulting mixture was allowed to incubate for three minutes. During this incubation period, 400
15 microliters of Solution A was added to the reaction medium of the immunoassay device prepared in Example 1 and allowed to drain through the reaction medium. The mouse anti-BE IgG/saliva mixture was then added to the reaction medium and allowed to incubate for
20 two minutes. Another 400 microliter aliquot of Solution A was then added to the reaction medium and allowed to drain. After the Solution A had finished draining, a finger of the person who had provided the saliva sample was painted with 15 microliters of
25 a dilute solution of colloidal gold conjugated goat anti-mouse immunoglobulin G in a TRIS buffer containing 1.0% BSA, 0.05% Tween-20 and 0.05% azide (hereinafter "Gold Label Solution"). The finger was gently pressed against the reaction medium of the
30 immunoassay device and held in place for three seconds. The finger was then carefully rolled off the reaction medium. The reaction medium was

allowed to incubate for fifteen seconds before another 400 microliter aliquot of Solution A was added to the reaction medium. After completion of the procedure, the reaction zone was colored, showing a negative test result for benzoylecgonine (e.g., as shown with reaction zones 32-34 in Figure 2).

The determination was repeated except that a minor amount of benzoylecgonine was added to the saliva sample just prior to the addition of the mouse anti-BE IgG solution. In this instance, after completion of the test, the reaction zones were completely white (e.g., as shown with reaction zones 35-37 in Figure 2). At the completion of both determinations, the central well control zone contained an imprint of the finger of the person who had provided the saliva sample (e.g., as shown with the fingerprint 62 on control zone 16 in Figure 2).

EXAMPLE 3.

An immunoassay device is prepared according to the procedure of Example 1 except that the device contains six reaction zones for different analytes. Each reaction zone contains polystyrene particles coated with a different analyte conjugate embedded in the membrane. Reaction zones 32-37 contain polystyrene latexes coated respectively with cocaine (32), opiates (33), PCP (34), amphetamine/methamphetamine (35), tetrahydrocannabinol (36) and alcohol (37) as the analytes (as shown in Figure 1). The central control zone 16 is prepared as in Example 1 with polystyrene particles, coated with goat anti-mouse immunoglobulin G, embedded in the

membrane.

5 A saliva sample is obtained from a person to be tested for the presence or absence of the six analytes. A 200 microliter aliquot of a dilute solution of mouse anti-analyte immunoglobulin G for each of the six different analytes in a phosphate buffered saline (hereinafter "mouse anti-analyte IgG Solution I") is added to 200 microliters of the saliva sample. As described in Example 2, while 10 this mixture is incubating, the reaction medium is pretreated with Solution A. After the mouse anti-analyte IgG Solution I/saliva mixture is added to the reaction medium, the remainder of the test procedure described in Example 2 is followed. The 15 reaction medium is treated with Solution A, contacted with a finger of the test subject that has been coated with a dilute solution of colloidal gold conjugated goat anti-mouse immunoglobulin G and then treated again with Solution A. After this procedure 20 is carried out, central control zone 16 contains the imprint of the finger of the person providing the urine sample. Reaction zones 32-37 are all colored, showing a negative test result for the presence of all six analytes.

25 The remaining saliva is then spiked with minor amounts of amphetamine/methamphetamine, tetrahydrocannabinol and alcohol. A 200 microliter aliquot of mouse anti-analyte IgG Solution I is then added to 200 microliters of the spiked saliva and 30 the determination is repeated as described for the unspiked sample. After the procedure is carried out, the assay device exhibits the result shown in Figure 2. Central control zone 16 contains the

imprint of the finger of the person providing the urine sample. Reaction zones 32-34 show a negative test result and are colored. Reaction zones 35-37 show a positive result and are white.

5 EXAMPLE 4.

 An immunoassay device, containing six reaction zones, is prepared according to the procedure of Example 1. As in Example 3, reaction zones 32-37 respectively contain polystyrene latexes coated with cocaine (32), opiates (33), PCP (34), amphetamine/
10 methamphetamine (35), tetrahydrocannabinol (36) and alcohol (37) as the analytes. In this instance, however, the central control zone 16, prepared as in Example 1, contains polystyrene particles coated
15 with mouse anti-goat immunoglobulin G embedded in the membrane. A urine sample is obtained from a person to be tested for the presence or absence of the six analytes. A 200 microliter aliquot of the urine is mixed with 200 microliters of mouse
20 anti-analyte IgG Solution I and the test procedure is then carried out exactly as described in Example 2. After completion of the determination, the central control zone 16 contains the imprint of the finger of the person providing the saliva sample and
25 reaction zones 32-37 are all colored, showing a negative test result for the presence of all six analytes.

 The remaining urine is then spiked with minor amounts of amphetamine/methamphetamine,
30 tetrahydrocannabinol and alcohol. A 200 microliter aliquot of mouse anti-analyte IgG Solution I is then added to 200 microliters of the spiked urine and the

determination is repeated as described for the unspiked sample. After completion of the determination, the immunoassay device exhibits the result shown in Figure 2. Central control zone 16 contains the imprint of the finger of the person providing the saliva sample. Reaction zones 32-34 are colored (negative result) and reaction zones 35-37 are white (positive result).

EXAMPLE 5.

The reaction medium 83 of the immunoassay device shown in Figures 9 and 10 is prepared according to the procedure described in Example 1. The reaction medium 83 has six reaction zones 32-37, which contain polystyrene latexes coated respectively with cocaine (32), opiates (33), PCP (34), methamphetamine (35), tetrahydrocannabinol (36) and alcohol (37) as the analytes and a central control zone 16 containing polystyrene particles, coated with mouse anti-goat immunoglobulin G, embedded in the membrane (as shown in Figures 1).

A sterile swab 85 is placed under the tongue of a person whose saliva is to be tested for the presence or absence of the six analytes. After one minute, the swab 85 is removed and is placed in the incubation channel 80 and basin 81 of the immunoassay device (as shown in Figure 9). Three drops of a dilute solution of silver coated gold microsome conjugates of mouse anti-analyte immunoglobulin G for each of the six different analytes (hereinafter "mouse anti-analyte IgG Solution II") are added to the saliva swab. The treated swab is allowed to incubate for one minute

before a finger 82 of the person providing the saliva sample is pressed against the swab for ten seconds. The finger is then lifted off the swab, immediately pressed on the reaction medium of the immunoassay device and held in place for thirty seconds. The finger is then lifted off the reaction medium and the medium is allowed to incubate for two minutes before the result is read. At this, point the central control zone 16 contains the imprint of the finger of the person providing the saliva sample and reaction zones 32-37 are all colored, showing a negative test result for the presence of all six analytes.

The test procedure is repeated using a new sterile swab except that prior to the addition of the mouse anti-analyte IgG Solution II to the swab, a drop of a solution containing minor amounts of methamphetamine, tetrahydrocannabinol and alcohol is added to the swab. After completion of the test, the immunoassay device exhibits the result shown in Figure 2. Central control zone 16 contains the imprint of the finger of the person providing the saliva sample. Reaction zones 32-34 are colored (negative result) and reaction zones 35-37 are white (positive result).

EXAMPLE 6.

An immunoassay device is prepared as in Example 5. A saliva sample is collected from under the tongue of a person to be tested for the presence or absence of the six analytes using a sterile swab. While the saliva sample is being collected, 800 microliters of mouse anti-analyte IgG Solution I is

placed in a sample tube together with one drop of a dilute solution of PCP. The saliva-containing swab is allowed to soak in the solution in the sample tube for two minutes. A 400 microliter aliquot of the resulting mixture is applied to the reaction medium and allowed to incubate for twenty seconds. A 400 microliter aliquot of Solution A was then added to the reaction medium and allowed to drain. After the Solution A had finished draining, a finger of the person who had provided the saliva sample is painted with 15 microliters of Gold Label Solution. The finger is immediately pressed gently against the reaction medium of the immunoassay device and held in place for five seconds. After the finger is carefully rolled off the reaction medium, the medium is allowed to incubate for fifteen seconds before another 400 microliter aliquot of Solution A is added to the reaction medium. After the completion of the procedure, the reaction zone is completely white, showing a positive test result for PCP. The central control zone contains an imprint of the finger of the person who had provided the saliva sample.

EXAMPLE 7.

An immunoassay device is prepared according to the procedure of Example 1. The reaction medium comprises a nylon membrane. The reaction zone includes polystyrene particles coated with human serum albumin-cocaine embedded in the membrane. The central control zone includes polystyrene particles, coated with goat anti-human immunoglobulin G, embedded in the membrane.

A finger of a person to be tested for the presence or absence of cocaine in their blood is pricked with a needle to obtain a droplet of blood. The droplet is spread with a sterile spatula to form
5 a thin film of blood over the fingertip. The blood covered fingertip is pressed against the reaction medium and held in place for about ten seconds. After the finger is lifted off, the reaction medium is treated with a dilute solution of household
10 detergent, thereby removing the red blood color from the reaction medium. After the red blood color has been removed, the reaction medium is treated with a dilute solution of alkaline phosphatase conjugated mouse anti-cocaine immunoglobulin G and allowed to
15 incubate for two minutes. The reaction medium is then treated sequentially with Solution A, a dilute solution of 5-bromo-4-chloro-3-indolyl phosphate and nitroblue tetrazolium (BCIP/NBT) and again with Solution A. The reaction medium is allowed to
20 incubate for several minutes between the BCIP/NBT treatment and the final Solution A treatment. When the procedure is completed, the central control zone contained an imprint of the finger of the person who had provided the blood sample and the reaction zone
25 was colored, showing a negative test result for cocaine.

EXAMPLE 8.

An immunoassay device is prepared as in Example 7. A 500 microliter aliquot of a dilute solution of
30 mouse anti-cocaine immunoglobulin G in a phosphate buffered saline containing 0.1% bovine serum albumin (BSA) and 0.05% Tween-20 (hereinafter "mouse

anti-cocaine IgG Solution"), is placed in a small sample tube. One drop of a solution containing a minor amount of cocaine is added to the sample tube.

5 A finger of a person to be tested for the presence or absence of cocaine in their blood is pricked with a needle to obtain a droplet of blood and two droplets of blood are added to the mixture in the sample tube. After the resulting mixture is allowed to incubate for two minutes in the sample
10 tube, the mixture is added to the reaction medium of the immunoassay device. The mixture is incubated for one minute on the reaction medium, which is then rinsed with Solution A. A finger of the person who had provided the blood sample is painted with 15
15 microliters of Gold Label Solution. The finger is gently pressed against the reaction medium of the immunoassay device, held in place for five seconds and then carefully rolled off the reaction medium. The reaction medium is allowed to incubate for
20 fifteen seconds before another 400 microliter aliquot of Solution A is added to the reaction medium. After completion of the procedure, the reaction zone is completely white (positive test for cocaine) and the control zone contains the imprint
25 of the finger of the person providing the blood sample.

EXAMPLE 9.

An immunoassay device is prepared according to the procedure of Example 1 except that the device
30 contains six reaction zones (see Figure 1). Each reaction zone contains polystyrene particles coated with a different amount of cholesterol conjugate

embedded in the membrane. Reaction zone 32 contains a cholesterol conjugate which will produce a positive signal when contacted with a biological fluid having greater than or equal to 150 milligram/deciliter total cholesterol level. Reaction zones 33-36 contain intermediate amounts of cholesterol conjugate, wherein the amount of cholesterol conjugate increases sequentially from zone 32 to zone 35, i.e., 180 mg/dl, 240 mg/dl, 280 mg/dl. The reaction medium also includes two reference control zones corresponding to 30 ml/dl and 65 ml/dl high density lipoprotein.

A finger of a person to be tested for the amount of cholesterol in their blood is pricked with a needle to obtain a droplet of blood. The droplet is added to a small sample tube containing 400 microliters of a dilute solution of mouse anti-cholesterol immunoglobulin G and colloidal gold conjugated mouse anti-cholesterol immunoglobulin G in a phosphate buffered saline containing 0.1% bovine serum albumin (BSA) and 0.05% Tween-20 (hereinafter "mouse anti-cholesterol IgG Solution"). After incubation for three minutes, a thin film of the mixture is spread over a fingertip of the person providing the blood sample using a sterile spatula. The fingertip is immediately gently pressed against the reaction medium of the immunoassay device. After holding the finger in place for five seconds, the finger is carefully lifted off the reaction medium. The reaction medium is incubated for one minute and then treated with a dilute solution of household detergent to remove red blood color from the reaction medium. After completion of the

procedure, the control zone contains the imprint of the finger of the person providing the blood sample. Reaction zones 32-33 are colored (negative test for cholesterol), while reaction zones 34 and 35 are white (positive test for cholesterol). Each reaction zone gives a positive test for cholesterol when at least the predetermined threshold amount of cholesterol is present. The threshold cholesterol concentration differs for each reaction zone in correspondence to the amount of cholesterol conjugate adsorbed in the reaction medium in that zone.

EXAMPLE 10.

A listing of a number of other drugs, the presence or absence of which may be determined using the present invention follows. This is only exemplary and is not meant to limit the invention thereto.

<u>β-Blockers</u>	<u>Diuretics</u>
Acebutolol	Acetazolamide
Alprenolol	Amilonde
Atenolol	Bendroflumethiazide
Labetalol	Bumetanide
Metoprolol	Canrenone
Nadolol	Chlormerodrin
Oxprenolol	Chlorthalidone
Propanolol	Diclofenamide
Sotalol and related compounds	Ethacrynic acid
	Furosemide
	Hydrochlorothiazide
	Mersalyl
	Spironolactone
	Triamterene and related compounds

	<u>Stimulants</u>	<u>Narcotic analgesics</u>
	Amfepramone	Alphaprodine
	Amphetamine	Anileridine
5	Amphetaminil	Suprenorphine
	Amiphenazole	Codeine
	Benzphetamine	Dextromoramide
	Caffeine	Dextropropoxyphene
	Cathine	Diamorphine
10	Chlorphentermine	Dihydrocodeine
	Clobenzorex	Dipipanone
	Clorprenaline	Ethoheptazine
	Cocaine	Ethylmorphine
	Cropropamide	Levorphanol
	Crotethamide	Methadone
15	Dimethamphetamine	Morphine
	Ephedrine	Nalbuphine
	Etafedrine	Pentazocine
	Ethamivan	Pethidine
20	Etilamphetamine	Phenazocine
	Fencamfamin	Trimeperidine and related compounds
	Fenethylline	
	Fenproporex	
	Furfenorex	
	Mefenorex	
25	Methamphetamine	
	Methoxyphenamine	
	Methylephedrine	
	Methylphenidate	
	Morazone	
30	Nikethamide	
	Pemoline	
	Pentetrazol	
	Phendimetrazine	
	Phenmetrazine	
35	Phentermine	
	Phenylpropanolamine	
	Pipradrol	
	Prolintane	
	Propylhexedrine	
40	Pyrovalerone	
	Strychnine and related compounds	

5	<u>Anabolic Steroids</u> Bolasterone Boldenone Clostebol Dehydromethyltestosterone Fluoxymesterone Mesterolone Methandienone Methandrostenolone Methenolone Methyltestosterone Nandrolone Norethandrolone Oxandrolone Oxymesterone Oxymetholone Stanozolol Testosterone and related compounds	<u>Hallucinogens</u> Lysergic Acid Diethylamide Mescaline Phencyclidine (PCP) Ketamine 2,5-Dimethoxy-4-Methylamphetamine Tetrahydrocannabinol Marijuana
20	<u>Opiates</u> Heroin Morphine Methandone Meperidine Codeine Propoxyphene	<u>Sedatives/Hypnotics</u> Chloral Hydrate Glutethimide Meprobamate Methaqualone
30	<u>Barbiturates</u> Amobarbital Pentobarbital Secobarbital Phenobarbital Butalbital Butabartial	<u>Benzodiazepines</u> Diazepam Clorazepate Chlordiazepoxide Oxazepam Flurazepam Lorazepam Alprazolam
35	<u>Antipsychotics/ Antidepressants</u> Chlorpromazine Trazodone Haloperidol Amoxapine Lithium Carbonate Imipramine	<u>Solvents</u> Ethanol Methanol Isopropanol Ethylene Glycol Chloroform

5	<u>Analgesics</u> Acetylsalicylic Acid Acetaminophen Ibuprofen Diflunisal Phenylbutazone	<u>Anabolic Steroids</u> Testosterone Methyltestosterone Nandrolone Stanozolol Oxandrolone Methandrostenolone Clostebol Mesterolone Norethandrolone
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EXAMPLE 11.

The following table sets forth exemplary chromogenic substrates yielding water-insoluble products that may be used with an appropriate enzyme conjugate in the invention, instead of the colloidal gold label previously noted. Labeling methods utilizing enzyme/chromogen couples are well known and would be easily practiced by one skilled in the art.

Chromogenic Substrates Yielding
Water-Insoluble Products

	<u>Enzyme</u>	<u>Substrate</u>	<u>Abbrev- iation</u>	<u>Orig. Color</u>	<u>Final Color</u>
5	Horseradish Peroxidase	Diaminobenzidene	DAB	Clear	Brown
		Diaminobenzidene with nickel enhancement	DAB/ nickel	Clear	Grey/ Black
10		3-Amino-9- ethylcarbazole	AEC	Clear	Red
		4-Chloro-1- naphthol	---	Clear	Blue
15	Alkaline Phosphatase	Naphthol-AS-BI- phosphate/fast red TR	NABP/ FR	Clear	Red
		Naphthol-AS-MX- phosphate/fast red TR	NAMP/ FR	Clear	Red
20		Naphthol-AS-BI- phosphate/new fuchsin	NABP/ NF	Clear	Red
	Purple	Bromochloroindolyl		BCIP/	Clear
25		phosphate/nitro- blue tetrazolium	NBT		
		5-Bromo-4-chloro- 3-indolyl-B-d- galactopyranoside	BCIG	Clear	Blue
30	B-Galacto- sidase	Naphthol AS-BI-B- d-galactopyrano- side	NABG	Clear	Red

EXAMPLE 12.

In this example, the device and method of Example 7 were used, except that the assay device included reaction zones for the analytes, shown in
5 Example 3, the surface of the reaction medium was pre-wetted with Solution A, the test subject's finger was pricked to produce a sufficient quantity of blood to smear the blood over the tip of the subject's finger, and the smeared blood was allowed
10 to dry on the subject's finger. The finger with the dried blood was then applied to the surface of assay device and held in contact with the device for about 15 seconds. After the finger is lifted off, the reaction medium was treated with a dilute solution
15 of clearing reagent, thereby removing the red blood color from the reaction medium. After the red blood color was removed, the reaction medium is treated with a dilute solution of conjugates of colloidal gold bonded to a secondary antibody, each conjugate
20 specifically binding to one of the named analytes. After incubation for about two minutes, a fingertip which had not been used to apply the blood was coated with about 10 microliters of colloidal gold conjugated to a ligand specific for a ligand
25 receptor in the control zone. After the fingertip was removed, the central control zone contained an imprint of the finger of the person who had provided the blood sample and the reaction zone was colored, showing a negative test result for the analytes.

30 EXAMPLE 13.

In this example, the device and method of Example 7 were used, except that the assay device included reaction zones for the analytes, shown in

Example 3, the surface of the reaction medium was dry, the test subject's finger was pricked to produce a sufficient quantity of blood to smear the blood over the tip of the subject's finger, and the smeared blood was immediately (before drying) applied to the reaction medium for about 15 seconds. After the finger is lifted off, the reaction medium was allowed to dry (typically about 1 to 2 minutes), treated with a clearing solution to remove the red blood cells, and treated with a dilute solution of clearing reagent, thereby removing the red blood color from the reaction medium. After the red blood color was removed, the reaction medium is treated with a dilute solution of conjugates of colloidal gold bonded to a secondary antibody, each conjugate specifically binding to one of the named analytes. After incubation for about two minutes, a fingertip which had not been used to apply the blood was coated with about 10 microliters of colloidal gold conjugated to a ligand specific for a ligand receptor in the control zone. After the fingertip was removed, the central control zone contained an imprint of the finger of the person who had provided the blood sample and the reaction zone was colored, showing a negative test result for the analytes.

EXAMPLE 14.

An assay device, containing five reaction zones (corresponding to areas 32 and 34-37 in Figure 1), is prepared according to the procedure of Example 1. As in Example 3, the reaction zones contain polystyrene latexes coated respectively with cocaine (32), PCP (34), amphetamine/ methamphetamine (35), tetrahydrocannabinol (36) and alcohol (37) as the

analytes. In this instance, however, zone 33 does not contain any adsorbed analyte or antibody and serves as a reference control zone, i.e., a zone which will remain completely white at the conclusion of the test. The central control zone ("identity control zone" corresponding to zone 16 in Figure 1) is prepared as in Example 1, except that it contains polystyrene particles coated with goat anti-rabbit immunoglobulin G embedded in the membrane.

5
10 The reaction medium is treated with 500 microliters of Solution A, thereby wetting the reaction medium. A sample of blood is obtained from a test subject to test for the presence or absence of the five analytes in their blood. A portion of
15 the sample is coated on the test subject's fingertip and applied to the five reaction zones and the reference control zone. Pressure is maintained on the reaction medium for about 10-15 seconds.

A few drops, e.g., about a 500 microliter
20 aliquot of a dilute household detergent solution is then added to the reaction medium and allowed to drain. The household detergent treatment lyses any red blood cells present in the reaction medium and rinses the dark red color from the blood out of the
25 medium. The five reaction zones and the reference control zone are then treated with about 10 ml of a dilute solution of a mixture of colloidal gold conjugated mouse anti-analyte immunoglobulin G in a TRIS buffer containing 1.0% BSA, 0.05% Tween-20 and
30 0.05% azide.

A finger of the person who had provided the blood sample is then painted with 15 microliters of a dilute solution of colloidal gold conjugated rabbit anti-goat immunoglobulin G in a TRIS buffer

containing 1.0% BSA, 0.05% Tween-20 and 0.05% azide. The finger is gently pressed against the identity control zone of the reaction medium and held in place for three seconds. After the finger is
5 carefully lifted off the reaction medium, the medium is allowed to incubate for fifteen seconds before another 400 microliter aliquot of Solution A is added to the reaction medium.

After completion of the procedure, the reaction
10 zones 32 and 34-37 are colored, showing a negative test result for the analytes. The reference control zone 33 is completely white and the identity control zone contains the imprint of the finger of the person providing the blood sample.

15 Although the present invention has been described in terms of exemplary embodiment, it is not limited to these embodiments. Alternative embodiments, examples, and modifications which would still be encompassed by the invention may be made by
20 those skilled in the art, particularly in light of the foregoing teachings. Therefore, the following claims are intended to cover any alternative embodiments, examples, modifications, or equivalents which may be included within the spirit and scope of
25 the invention as defined by the claims.

Claims:

1. A self-contained assay test device comprising:
 - (a) a housing;
 - 5 (b) a reaction medium having at least one reaction zone located in said housing;
 - (c) a medium containing a signal-producing agent located in said housing; and
 - (d) a medium for containing a test sample
 - 10 located in said housing, each of said media (c) and (d) being movable independently between a first position in spaced relationship to said reaction medium and a second position in substance-transferable contact with said reaction medium.
- 15 2. A self-contained assay test device according to claim 1 wherein said reaction medium also includes a control zone to identify a test subject.
- 20 3. A self-contained assay test device according to claim 1 wherein said at least one reaction zone includes a reagent for determining the presence or absence of analyte in a test sample.
- 25 4. A self-contained assay test device according to claim 3 wherein said analyte is a drug or drug metabolite.
5. A self-contained assay test device according to claim 1 wherein said reaction zone includes a drug or drug metabolite.

6. A self-contained assay test device according to claim 1 wherein said at least one reaction zone includes a member of a ligand/receptor pair.

5 7. A self-contained assay test device according to claim 6 wherein said member of a ligand/receptor pair is a member of an antigen/antibody pair.

10 8. A self-contained assay test device according to claim 1 wherein said reaction medium is a low protein-binding, hydrophilic polysulfone membrane.

15 9. A self-contained assay test device according to claim 2 wherein said control zone includes a member of a ligand/receptor pair.

10. A self-contained assay test device according to claim 1 wherein said reaction medium includes a member of a ligand/receptor pair which forms a conjugate with a human antibody to a drug.

20 11. A self-contained assay test device according to claim 1 wherein said signal-producing agent is bound to a member of a ligand/receptor pair.

25 12. A self-contained assay test device according to claim 1 wherein said signal-producing agent is a member of a ligand/receptor pair labelled with colloidal gold.

13. A self-contained assay test device according to claim 9 wherein said signal-producing agent forms a conjugate with said member of a ligand/receptor pair in said control zone.

5 14. A self-contained assay test device according to claim 1 wherein said medium for containing a test sample includes a test reagent capable of reacting with an analyte being sought.

10 15. A self-contained assay test device according to claim 1 wherein said medium for containing a test sample includes a member of a ligand/receptor pair capable of forming a conjugate with an analyte being sought.

15 16. A self-contained assay test device according to claim 1 wherein said medium for containing a test sample is mounted on one end of a cylinder.

20 17. A self-contained assay test device according to claim 1 wherein said medium for containing a test sample comprises a hydrophilic absorbent medium.

25 18. A self-contained assay test device according to claim 17 wherein said medium for containing a test sample comprises a hydrophilic sponge material.

19. A self-contained assay test device according to claim 1 wherein substance transferable contact occurs at a pressure of 2 to 15 psi.

20. A self-contained assay test device comprising:

a two-part housing;

a reaction medium having at least one reaction
5 zone mounted in a first part of said two-part housing;

a medium containing a signal-producing agent mounted in a second part of said two-part housing;

a removable medium for containing a test sample
10 located intermediate said first and second parts of said housing and in substance-transferable contact with said reaction medium when said first and second parts of said housing are in a first closed position of said housing; and

15 said medium containing a signal-producing agent being in substance transferable contact with said reaction medium when said first and second parts of second housing are in a first closed position .

21. A self-contained assay test device
20 according to claim 20 wherein said first and second parts of said housing are joined by a hinge.

22. A self-contained assay test device according to claim 20 wherein said reaction medium further includes a control zone having a member of
25 ligand/receptor pair for identification of a test subject and said at least one reaction zone includes a member of a ligand/receptor pair for determining the presence of an analyte in the test sample, said medium containing a signal-producing agent is a
30 member of a ligand/receptor pair labelled with colloidal gold, said removable medium for containing a test sample is a hydrophilic sponge material which

includes a member of a ligand/receptor pair capable of forming a conjugate with an analyte being sought and is mounted on one end of a cylinder.

23. An assay device comprising:

- 5 (a) a housing;
(b) a reaction medium having at least one reaction zone located in said housing; and
(c) a medium for containing a test sample located in said housing and movable between a first
10 position in spaced relationship to said reaction medium and a second position in substance-transferable contact with said reaction medium.

24. An assay device according to claim 23 wherein substance transferable contact occurs at a
15 pressure greater than atmospheric pressure.

25. An assay device according to claim 23 wherein substance transferable contact occurs at a pressure of 2 to 15 psi.

26. An assay device comprising:

- 20 (a) a housing;
(b) a reaction medium having at least one reaction zone located in said housing; and
(c) a medium containing a signal-producing agent located in said housing and movable between a
25 first position in spaced relationship to said reaction medium and a second position in substance-transferable contact with said reaction medium.

27. An assay test device according to claim 26 wherein said reaction medium also includes a control

zone to identify a test subject.

28. An assay test device comprising:

- (a) a housing;
- (b) a reaction medium having at least one
5 reaction zone located in said housing; and
- (c) a medium for containing a test sample
located in said housing and movable between a first
position in spaced relationship to said reaction
medium and a second position in substance-
10 transferrable contact with said reaction medium.

29. An assay test device according to claim 28
wherein said reaction medium also includes as
control zone to identify a test subject.

30. A kit for performing an assay comprising:

- 15 (i) an assay test device including:
 - (a) a housing;
 - (b) a reaction medium having at least one
reaction zone located in said
housing; and
 - 20 (c) a medium containing a signal-
producing agent located in said
housing and movable between a first
position in spaced relationship to
said reaction medium and a second
25 position in substance-transferable
contact with said reaction medium.
- (ii) a reagent including a member of a
ligand/receptor pair capable of forming a conjugate
with an analyte.

31. An assay kit according to claim 30 wherein said reaction medium also includes a control zone to identify a test subject.

32. An assay kit comprising:

- 5 (i) an assay test device including:
- (a) a housing;
 - (b) a reaction medium having at least one reaction zone located in said housing; and
 - 10 (c) a medium for containing a test sample located in said housing and movable between a first position in spaced relationship to said reaction medium and a second position in substance-
 - 15 transferable contact with said reaction medium.
- (ii) a signal-producing reagent capable of reacting with a member of a ligand/receptor pair.

33. A test kit according to claim 32 wherein
20 said reaction medium includes a control zone to identify a test subject.

34. A method for determining the presence of an analyte in a test subject comprising:

- (a) contacting a medium for containing and
25 transferring a test sample with a portion of a test sample when said medium for containing and transferring a test sample is in a first position spaced apart from a reaction medium having at least one reaction zone;
- 30 (b) moving said medium for containing and transferring a test sample to a second position in

substance-transferable contact with said reaction medium; and

5 (c) moving a medium containing a signal-producing agent from a first position in spaced relationship to said reaction medium to a second position in substance-transferable contact with said reaction medium; and

(d) determining the presence or absence of the analyte in the test sample from the reaction medium.

10 35. A method of determining the presence of an analyte according to claim 34 wherein said reaction medium includes a control zone and a part of the body of a test subject is placed in contact with the control zone after substance-transferable contact
15 occurs between the reaction medium and the medium containing a signal-producing agent.

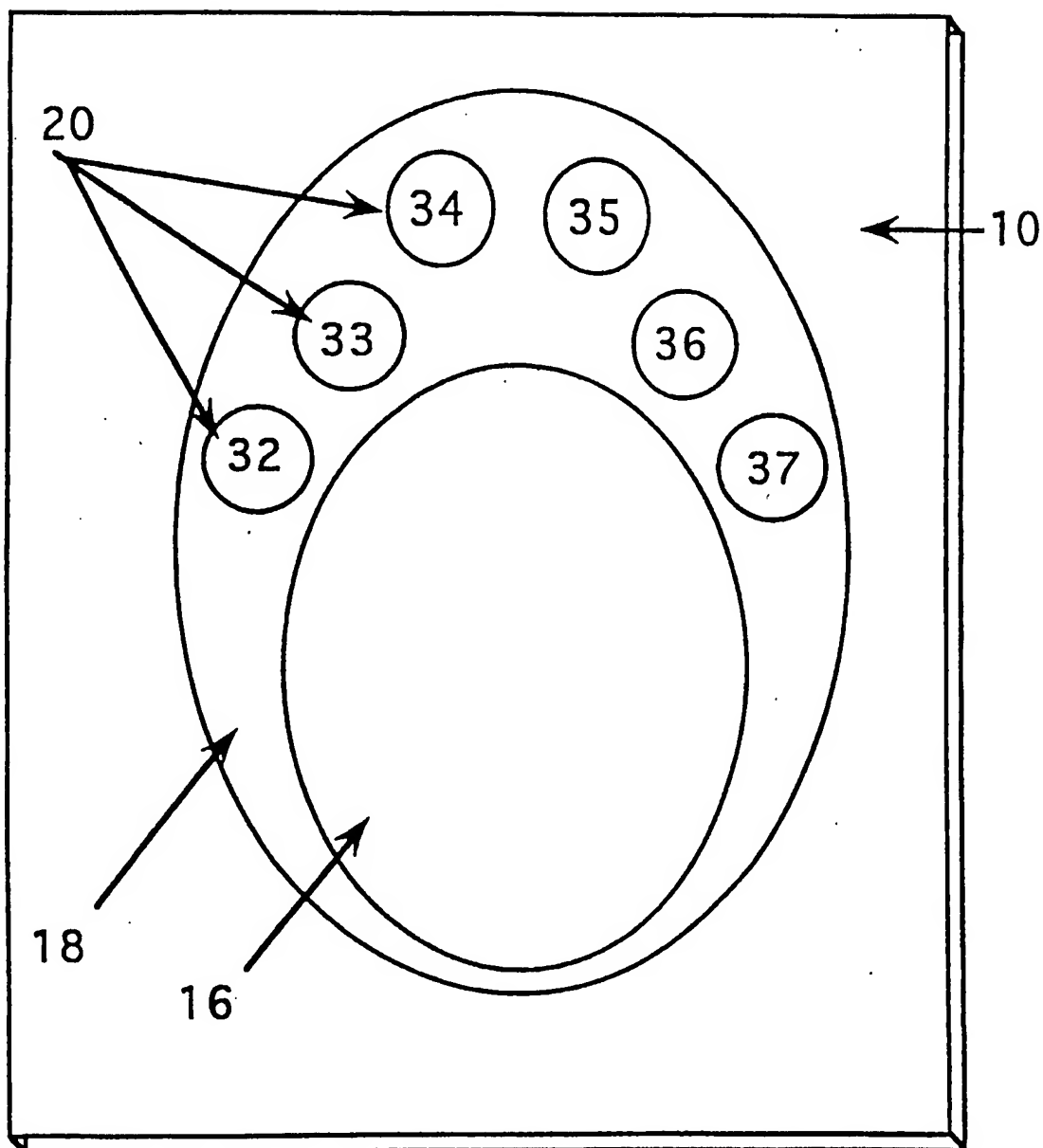


FIG. 1

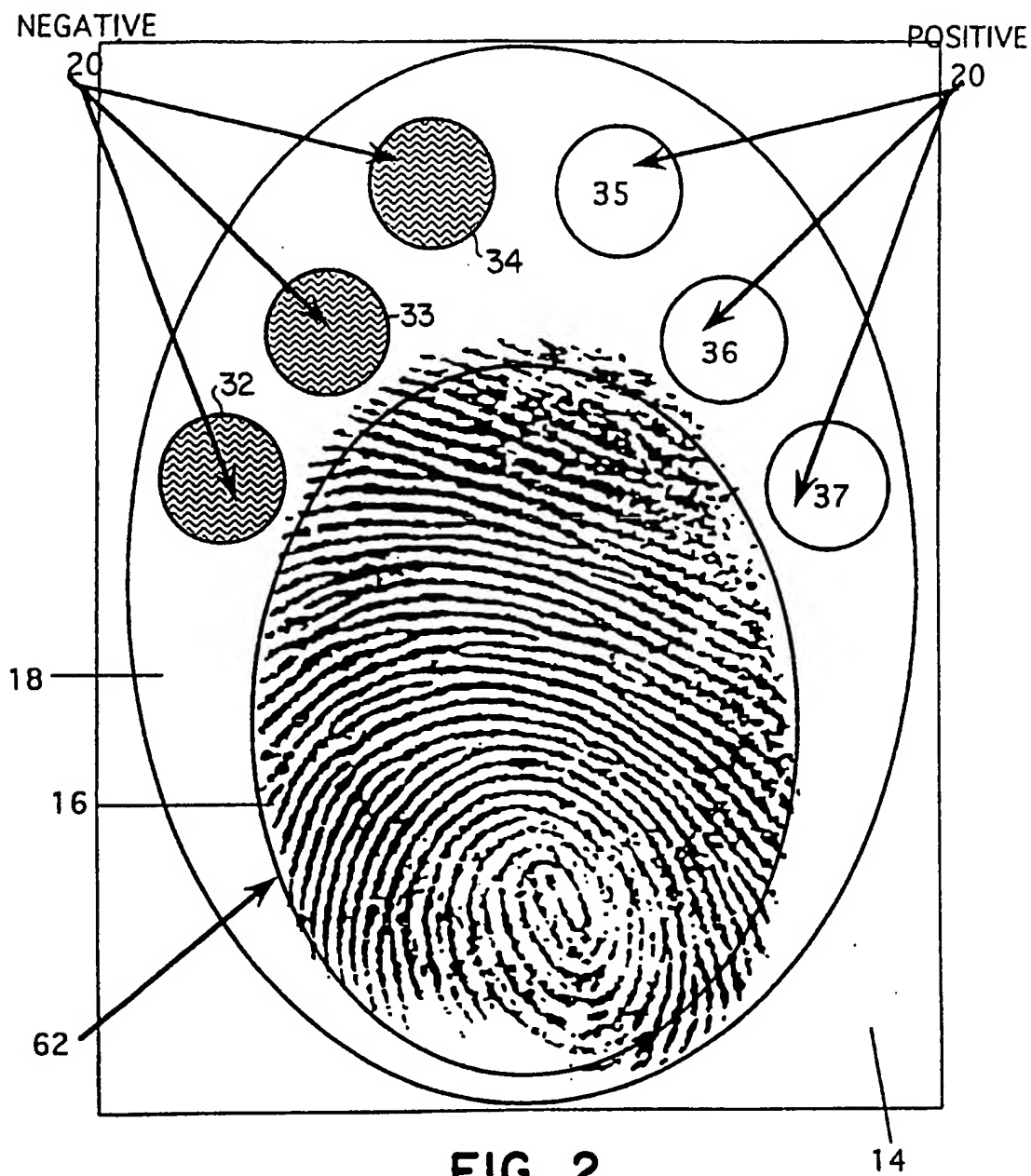


FIG. 2

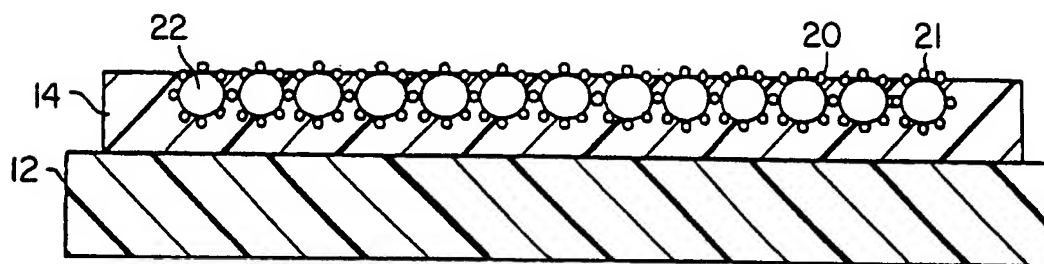


FIG. 3

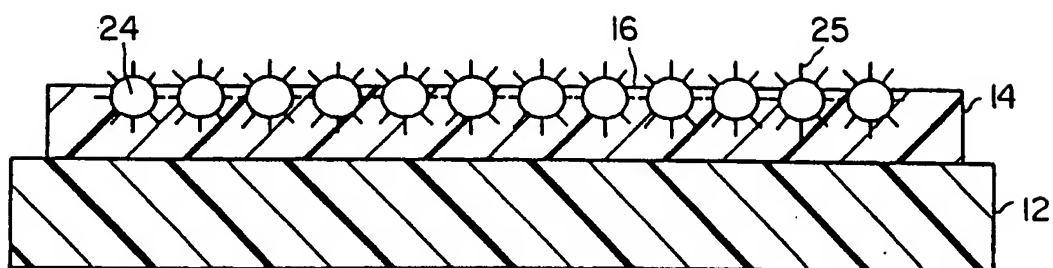


FIG. 4

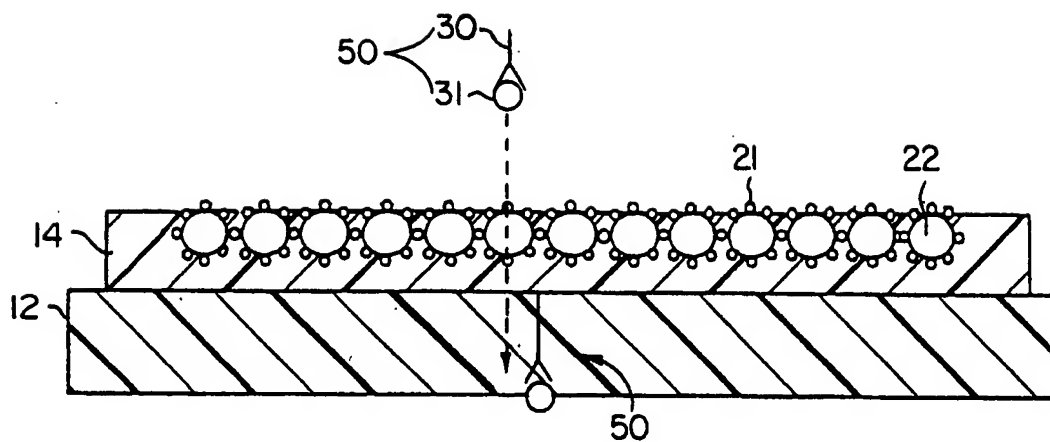


FIG. 5

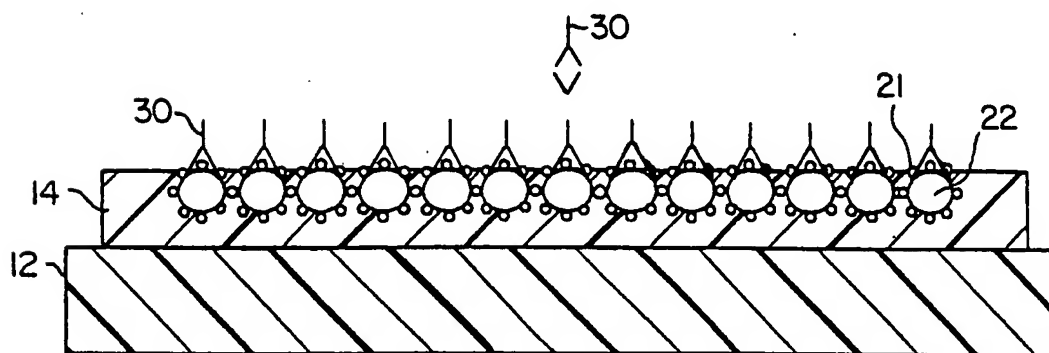


FIG. 6

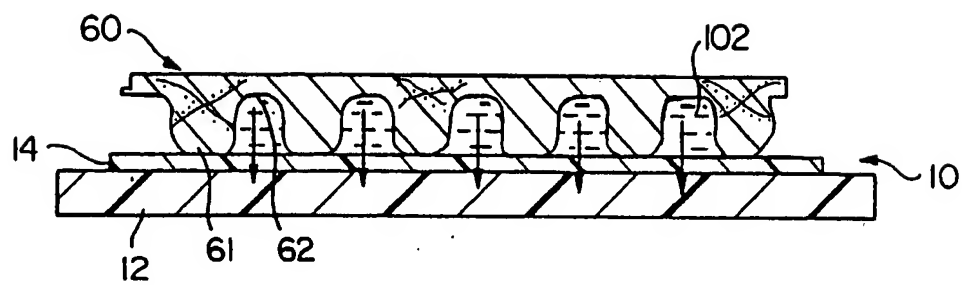


FIG. 7

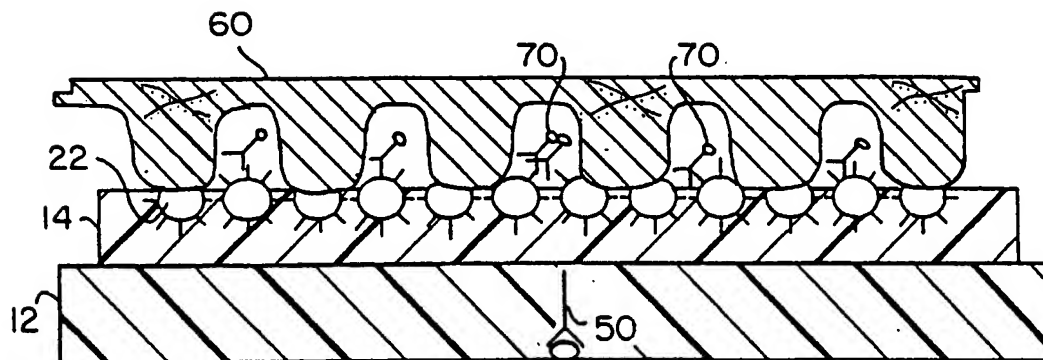


FIG. 8

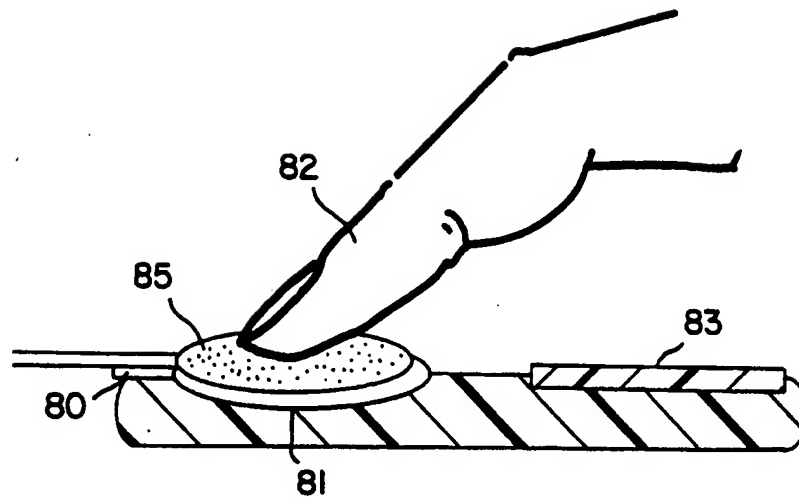


FIG. 9

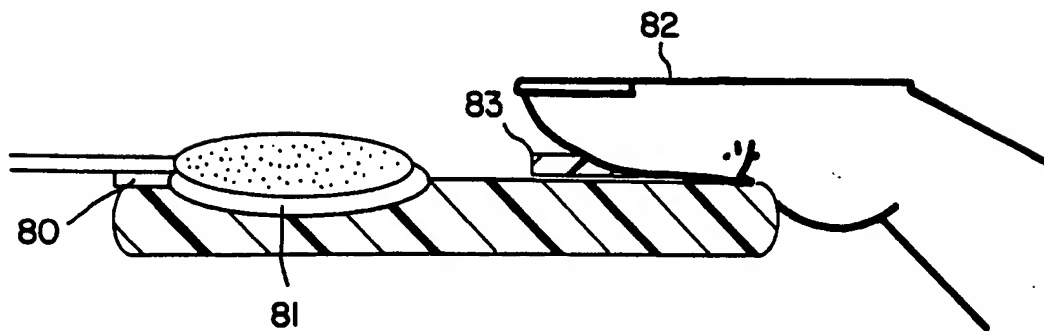
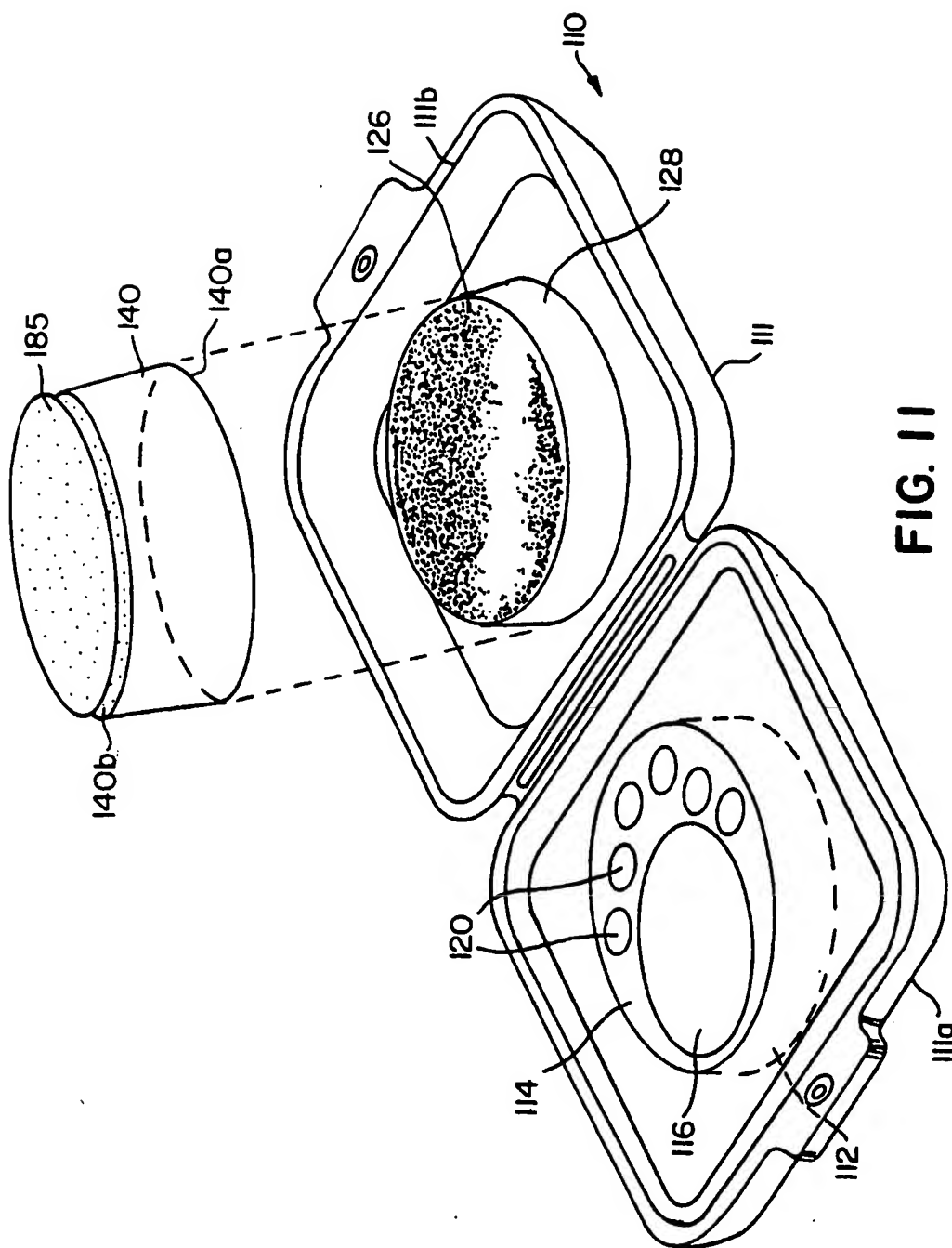


FIG. 10



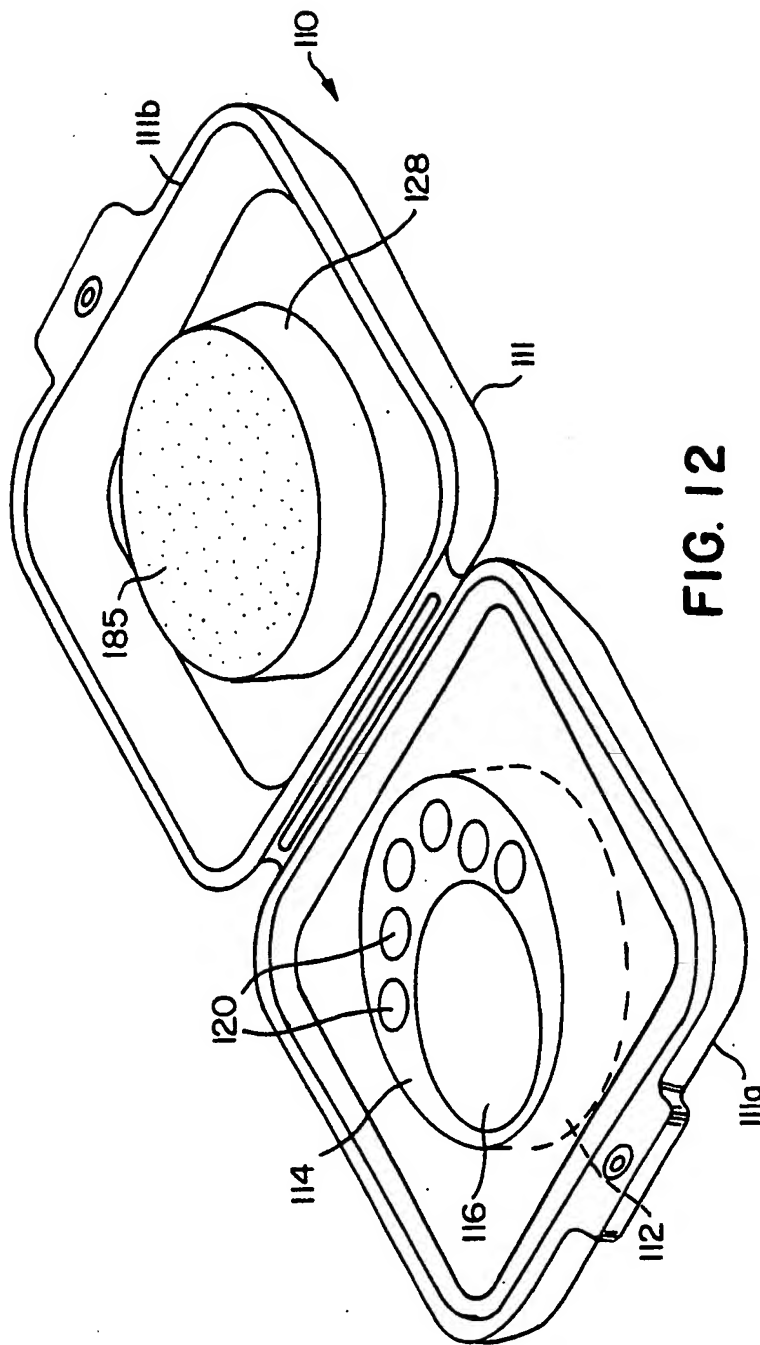
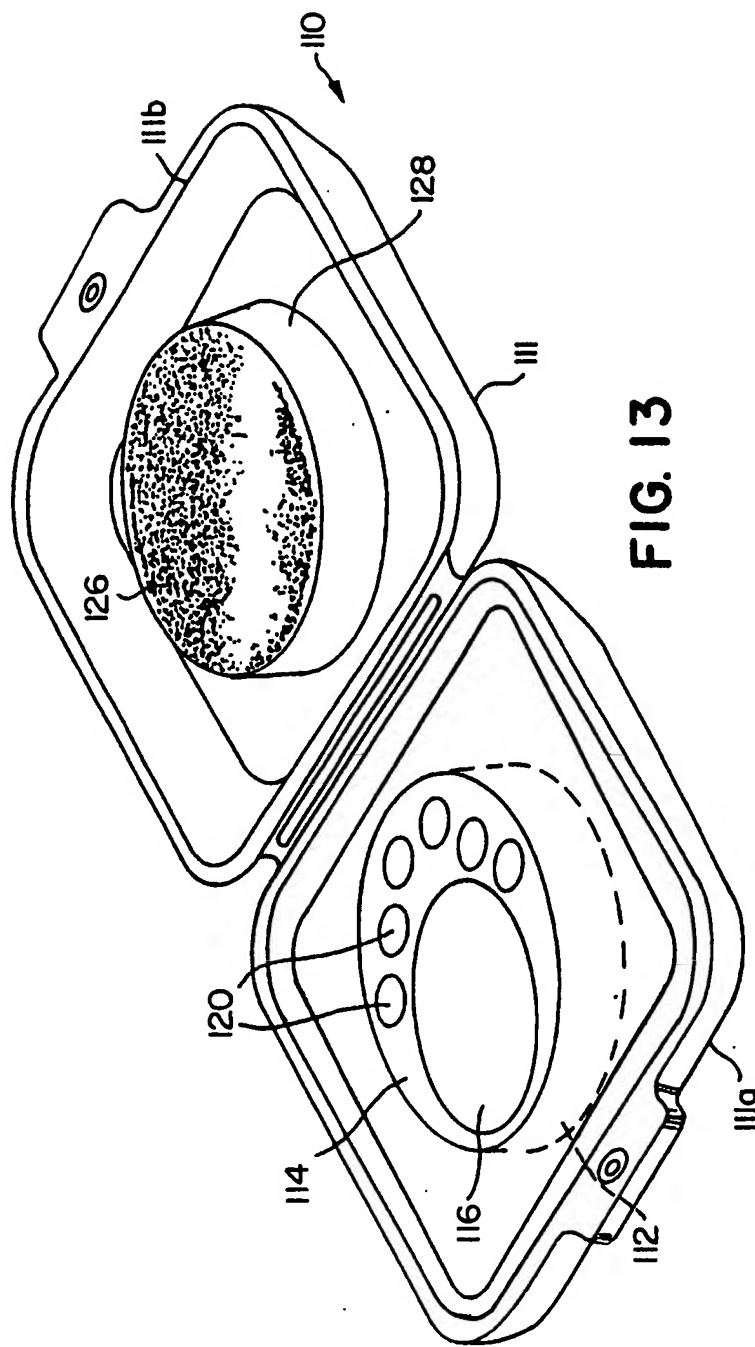


FIG. 12



INTERNATIONAL SEARCH REPORT

International application No.

PCT/US94/10473

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :G01N 33/53

US CL :436/534; 422/56

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 436/518, 525, 528, 529, 530, 531, 534, 808,810, 815, 901; 422/56, 58, 61

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
N/A

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS, MEDLINE, CAS, EMBASE, TOXICOL, WORLD PATENT INDEX

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X --- Y	US, A, 4,717,656 (SWANLJUNG) 05 January 1988, see entire document, especially see col. 5, line 48 to col. 6, line 22, col. 8, line 27 to col. 9, line 5 and Figures 5a and 5b.	1-3, 6-7, 9, 11, 13-15, 17-18, 23, 26-33, 34-35 ----- 4-5, 8, 10, 12, 16, 19, 20-22, 24-25
Y	US, A, 4,963,325 (LENNON ET AL.) 16 October 1990, see entire document, especially see col. 1, lines 14-24, col. 2, lines 36-42 and 45-48 and col. 3, line 33 to col. 4, line 13.	4-5, 10, 12, 16, 19-22, 24-25

☒ Further documents are listed in the continuation of Box C.
 ☐ See patent family annex.

* Special categories of cited documents:	*T	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
A document defining the general state of the art which is not considered to be of particular relevance	*X*	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
E earlier document published on or after the international filing date	*Y*	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*Z*	document member of the same patent family
O document referring to an oral disclosure, use, exhibition or other means		
P document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

20 NOVEMBER 1994

Date of mailing of the international search report

09 JAN 1995

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INTERNATIONAL SEARCH REPORT

International application No.
PCT/US94/10473

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	THE FISHER CATALOG, published 1990 by Fisher Scientific (Pittsburgh, PA), see page 738 "Supor Modified Polysulfone Membranes".	8
Y	US, A, 5,244,815 (GUIRGUIS) 14 September 1993, see entire document, especially claim 18.	1-35
A	US, A, 3,784,358 (DRAKE) 08 January 1974, see entire document, especially col. 4, lines 47-54.	1-35
A,P	US, A, 5,308,580 (CLARK) 03 May 1994, see entire document.	1-35